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	AI913553, AI418464, AA299849, AA729787 AI547277 AC005005 and A CODOMO	AI350354, AI904299, AI9022503, D61534, T78554, AW183962, AI218626, AW304978, W74167, AI081779, AL022238, and AL137499	T63995, and AI669879.	AI554514, AW117613, AA401269, AA89442, AI923609, AA968468, AA779787, W56250, AI049682, AI342711, AI277156, AA424469, AA443257, AW023482, AI978942, AI611714,	AL132/1/1, and AB024689.	AA683227, H65257, Z99396, AW392670, AW372827, AL119497, AL134531, U46341, AW384394, AL119457, AL119483, AL119418, AL119324, AL119396, AL038837, AL119484, AL119391, U46350, AL119341, AL119443, U46349,	AL03/031, AL036/25, U46351, AL119335, AL119363, AL119355, AL119496, AL036418, AA631969, U46347, AL043029, AL042989, AL119401, AL134920, AL134533, AL036034	AL042970, AL119488, AL119439, AL042965, AL134528, AI142139, AL042975, AL043033,	AL119444, AL043011, AL037205, AL042614, AL036858, AL134538, AL042984, AL042544,	AL042542, AL043019, U46346, AL043003, AL043037, AL042551, AL042896, AL042850,	AL042995, U46345, AL037094, AL036268, AL036190, AL037639, AL036196, AL037526,	ALU3/V63, ALU30998, ALU3/V62, ALU3844, ALU3/615, ALU36733, AR066494, AR060234, A81671, AR023813, AB026436, AR064707, AR069079, and AR064110	, ATTLANTING (1) (1) (1) (1) (1) (1) (1) (1) (1) (1)	AI678080, AW118890, AI962969, AI936455, AL037402, AA969064, AI127304, AI677882	AW195711, H19013, N33206, AI127565, AI332958, AI089991, AI253076, AW051831,	AI025225, AI004194, AI873265, AW182432, N48995, AA528395, AI290549, AI003987,	AA022465, AI377431, H81787, AA225942, N44742, H19307, W25684, AI246674, N95594,	H81/86, Z42919, R52370, H98158, A1382954, Z43457, A1244671, A1093458, A1042622,	K52369, R73912, W20350, Z39530, AA022998, C04474, W92421, R74008, F02397, N92731,	AA934795, AA502534, AW081759, H98461, AA744985, AA883984, AA319261, AI619688,	AL0/9491, N45/17, and AA662/87.	AL043767, AA043579, and H42377.	AW452627, and AC009248.	AI910578.					R90939, H66735, and R89394.
15 - 516	15 = 570	15 - 703	15 - 665	15 - 839		15 - 565							15 - 438	15 - 2645								15 - 274	15 - 362	15 - 273	15 - 352	15 - 441	15 - 265	15 - 403	15 - 490
1 - 502	1 - 556	1 - 689	1 - 651	1 - 825		1 - 551							1 - 424	1 - 2631								1 - 260	1 - 348	1 - 259	1 - 338	1 - 427	1 - 251	1 - 389	1 - 476
969161	961595	965306	954332	662596		969151							951658	961388				-			00.000	963100	574510	574232	574008	625096	932383	653242	531348
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HCOOM77	HCOOX10	HC00Z11	HCOPP18	нсоое11	7,1100011	НСООН12			•				нсоооол	HE8SG46							TIPAAAA	HEAAA42	HEAAB77	HEAAK46	HEAAM16	HEAAM52	HEAAN18	HEAAQ66	HEAAT36

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1 - 459	1 - 347	1 - 436	1 - 133	1 - 328	1 - 329				1 - 340	1 - 535										_												
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	1-312	1 - 293	1 - 333	1 - 335	1 - 496	1 - 430	1 - 372							•	-										
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									<u></u>						656299	676716	767284	823900	754344	965183			
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AA977410, T91051, AI828094, AI675983, AW248442, and AI681656	AI624142, AW393412, M62051, AI870044, AA3300RR and AB014538	AA707670.	AR064011.	AI813699, AW051520, AI312608, AI338779, AI369324, AI218925, R72618, R80387, R72688, R80403, A 4038076, D67386, A136827, BR0403, A186828, BR0403, BR0403, A186828, BR0403, A186828, BR0403, BR0	1173167 and 11900004 NOSGOS, RUZSGOS, ROSSGOS, and A1214650.	יייים מות ליייים אות ליייים אות היייים אות הייים אות היייים אות היייים אות היייים אות היייים אות היייים אות הייים אות היייים אות היייים אות היייים אות היייים אות היייים אות הייים את הייים הייים הייים את הייים היי	H83841, AA019657 and A1127286	R55547, H12557, H27260, H00825, R23780, AA747653, H12800, T50030, H42768, and	AW361580.	AL137660.	X66139, and X66140	AA335075 and AA 335406		AA335537, AA336174, AA335376, AW406878, A1541216, A1525220, A1525710	AL022318,	AA335399, and AA335663.	AA335412, AA335741 AA335606 and a a sorong	AA335523 and AA315948	AA335526 and AA347833	AA524859, AI366999, AW274554, AW085032, AA897153, AA334550, AA405462,	AW305246.	AA335591, and AA335639.	AA335593, AI247347, AA335892, AI700468. AI014987. AA629210 AI700878 and X66130	T31859, AI452722, AI810976, AA039492, AW166142, T34621, AA887990, AA 526699	AI491944, AI291744, AA971270, AI291429, AI147212, AI191377, AI282167, AW194181.	AI382209, AI819092, AI125991, AI291350, AA635803, AI076763, AI025483, AW054812,	AW026209, AI872247, AA446939, AA393844, AA595299, AA588205, AA635837, AA580350.	W42/14, AA652370, AI205639, AI346541, AI299347, AA618584, AA041546, AI190326,	AL389/81, AI038728, AA747482, AI374991, AI186987, AI272049, AA968514, AA781105,	AA/19399, AI/45517, AI479431, AW005070, AI760672, AI828575, AI653887, AI983727, AI650052, AI681964, A 4 6 5 3 6 8 9 6 9 6 9 6 9 6 9 6 9 6 9 6 9 6 9 6	H14119, AI588901, T67080, H20066, Z38557, AA069378, N95318, H19689, PR86071	AA069325, AW340398, AA532555, AA742707, AI095816, AI261987, T31864, AA224332.
	15 - 608	15-516	15 - 328	15 - 650	15 - 468	15 - 550	15 - 683	15 - 196		15 - 204	15 - 561	15 - 327	15 - 355	15 - 282		15 - 268	15 - 199	15 - 75	15-311	15 - 585		15 - 403	15 - 409	15 - 365								
	1 - 594	1 - 502	1 - 314	1 - 636	1 - 454	1-536	1 - 669	1 - 182		1 - 190	1 - 547	1-313	1-341	1 - 268		1 - 254	1 - 185	1 - 61	1 - 297	1-571		1 - 389	1 - 395	1 - 351								
	955291	887299	780837	934705	697419	706951	719387	906669		726316	950033	509456	557149	888726		509002	509452	586843	518331	925146		508706	508694	926914	-				-			
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	161	162	163	164	165	166	167	168	169										170	171	172	173	174	175	176	177	178	179	
	HEPAP12	HEPAS44	HEPBA39	HEPBB24	HEPBB60	HEPBG26	HEPBG35	HEPBH28	HEPBH38								-		HEPBH45	HEPBO69	HEPBQ47	нерво69	HEPBS10	HEPBX43	HEPCD36	HEPCE25	HEPCO59	HEPCT32	

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HEPCU32	HEQAE65	HEQAH70	HEQAO76	HEQAZ52	HEQBA41	HETAD29	HETAF20	HETAF49	HETAF89	HETAH16		HETAH66	HETAH67	HETAJ26	HETAK75													HETAN20	HETAP59	HETAP94	HETAR06	HETAR60		

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		918730	536192	509189	508655	720853	509099	523046	971505	999809	841924					921390	522829	525412			_	01100	883018	855500	cocce	028096		925489	208990	695021	525407
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		HETAT83	HETAZ13	HETBA01	HETBF45	HETBH48	HETBR25	HETBW39	HETCE12	HETCESS	HETCG63		•••			HETCH92	HETCL55	HETDA81				UETHN10	HETDE67	HETDE86	2077	HETDG67		HETDI03	HETDL92	HETDN90	HETDP21

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·	HETIG71	HETIJ84	HETIJ85	HETIQ89	HETIU60	HETIY84	HETJD30	HETJD86	·····	HETJG63	HETJI32	НЕТЖ67	HETJNSI	HETJT95	HETJX04	HETIY11						

	AUTOCOCON A ADDROGY AUTOCOCA A SECOND
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964389	944774	920690	746460
282	283	284	285
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	4		15 - 565	15 - 1289	15 - 509	
			1 - 551	1 - 1275	1 - 439	
			694216	947484	666273	
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7	791362	1 - 505	15 - 519	AA368122.
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	15 - 285	15 - 462	15 - 900		15 - 324	15 - 357	15 - 533	15 - 646		15 - 838	15 - 275	15 - 244	15 - 426	15 - 453	15 - 52	15 - 398	15 - 559	15 - 717	15 - 614	15 - 957			
·	1 - 271	1 - 448	1 - 332		1 - 310	1 - 343	1 - 519	1 - 632		1 - 824	1 - 261	1 - 530	1 - 412	1 - 439	1 - 38	1 - 384	1 - 545	1 - 703	1 - 600	1 - 943			
	925300	670215	577929		727699	926850	750779	002698	024071	147/18	708018	00/016	953283	868276	759843	973315	929647	856488	957804	914428			
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	HLWBS14	HLWBS43	HLWCA67		HLWCM44	HLWCM70	HLWCO66	HLWCQ53	6700/m Iri	HLWCQ62	HI WDA01	HI WDR18	HLWDD02	HLWDE60	HLWDL71	HLWEE76	HLWFG82	HLWFQ04	HMVDU41	HNBTP01			

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	15 - 242	15 - 341	15 - 483
	1 - 228	1 - 327	1 - 469
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	321	322	323
	HNBTT79	HNBTX52	e Mose

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	15 - 456	15 - 300	15 - 1595		15-1117
	1 - 442	1 - 286	1 - 1581		1 - 1103
	951814	958685	917723		917725
	324	325	326		328
	HNBUR07	HNGAO08	HNNNA06		HNNNA77

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	15 - 752	15 - 791	15 - 783	15 - 700
	1 - 738	1 - 777	1 - 769	1 - 686
	933730	969363	927903	964933
	329	330	331	332
	HNOAS06	HNOAX12	HNOBF57	HNOCQ04

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HNOC139	333	952611	1 - 101	15 - 115	AA806386, AI244963, AW069189, AA811454, AW069641, AA228016, AI926450, N20102, AA984503, AI283001, AI335808, A1022959, AI699076, AW008997, AI092767, AI358493, AI289652, AI309309, AA747446, AA521303, AI460092, AI354930, AW105294, AA902193, AI358182, AA975250, AA576719, AI766428, AI335889, AA071293, T10067, R58045, and AI355753.
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НОСМ 003	335	922418	1 - 602	15 - 616	
HOCPJ03	336	917484	1 - 299	15 - 313	AW374051, AW374048, AW374059, AA047322, AA573420, AA873293, H50745, T67156, and AI718619.
HODAD73	337	973463	1 - 490	15 - 504	AC004889,
HODAD95	338	974043	1 - 577	15 - 591	293783.
HODAG37	339	529410	1 - 224	15 - 238	
HODAH32	340	859509	1 - 305	15 - 319	AA076906, and AC004976.
HODAJ01	341	921666	1 - 570	15 - 584	
HODAJ35	342	529405	1 - 325	15 - 339	AC005618.
HODAK38	343	529404	1 - 310	15 - 324	
НОДАК95	344	960179	1 - 690	15 - 704	AI333350, AI522314, N93898, AI539488, AI276544, AW292555, AA194179, AA193323, AA621456, and N64007.
HODA016	345	529401	1 - 271	15 - 285	
HODAT56	346	529402	1 - 354.	15 - 368	
HODAV80	347	859519	1 - 287	15 - 301	
HODAW60	348	692684	1 - 315	15 - 329	
HODAW84	349	775425	1 - 503	15-517	AI800919, AI741507, N21056, AA966954 AA669758 AI018174 A1741507, N21056, AA966954
НОВВС01	350	651662	1 - 290	15 - 304	10075071, and A1045051.
HODBC07	351	954161	1-317	15 - 331	
НОВВЕ01	352	921655	1 - 102	15 - 116	
НООВН16	353	927781	1 - 286	15 - 300	
HODBO85	354	859559	1 - 156	15 - 170	
HODBT58	355	678444	1 - 303	15-317	AA224807, AI355986, AI791718, C14330, AA778962, N94325, AA312559, H71678, H19817

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	15 - 648	15 - 143	15-425	15-613	15 - 674	15 - 742	•	15 - 266	15 - 593	15 - 178	15 - 599	15 - 441	15 - 418	15 - 452	15 - 530	15 - 352	15 - 934		
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	HODEG86	НОБЕН08	HODEH22	HODEI13	HODEI48	HODEK50		HODEL92	HODEN75	HODEO87	HODEP04	HODEP12	HODEP86	НОБЕО28	HODEQ79	HODER91	HODES86		

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	421	422	423	424	425	426	427	428	429	430	431	432	433	434	435	436	437	438	439	440	441	442	443
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918456	859334	921529	859329	859314	921651	918637	934304																										
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	HODGO46	HODGP37	HODGP83	HODGP95	HODGO08	HODGQ22	НОДСО32	нордом	HODGT62	HODGW08	HODGX10	HODGX29	HODGZ06	HODGZ10	НОДНС03	HODHD16	HODHG71	НОДНН82	HODHI07	HODHI26	НОДНК02	HODHL 19	HODHL56	НОДНО11	HODHS34	HODHY53	HODJR03	HODJY33	HODJZ90	HODKN07	HOFAA15	HOFAB40	

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nor Age	800	766711	1 - 208	15 - 222	AI718967, A1066500, A1191408, AA970891, W16766, A1074219, F25014, N45385, AA465517, AA602119, N94010, F32882, F26209, F25830, N98860, A1934901, N77808, AA528414, A1338326, AW088372, AW102799, AA320204, AA768215, A1151192, AW419435, AA788658, A1150785, C03969, A1302844, AA725287, AL038871, A1350125, A1568272, AA983392, AA525200, AA977031, A1093521, A1580262, A1720008, H97078, A1313204, A1626036, AA526171, A8877563, A1144433, AA934574, A107769, AA573686, AA6739915, AA574008,
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nOraco	506	/0/359	1 - 372	15 - 386	AI694793, AW328387, AA287690, AW166132, AI701814, AI393309, AW005351, AI807923, AA304908, AA194090, AI799077, AI916382, AI131240, AI630546, AA308513, AW087574, AI138878, AA855025, AI188081, AW300307, AW340414, and R01958
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HOFMA24	511	782275	1 - 333	15 - 347	R06046, N24376, AA431932, H96741, AI632470, AW130380, AW197748, AI183539, and AB032969.
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HOFMB83	513	572839	1 - 343	15 - 357	
HOFMD13	514	916959	1 - 393	15 - 407	
HOFME41	515	867993	1 - 354	15 - 368	X52381, AF163863, M64428, X56219, AF076492, and X56223.
HOFMF03	216	924679	1 - 434	15 - 448	
HOFMF70	517	734917	1 - 374	15 - 388	AA078777, AW205164, M59936, AF099730, AJ004856, and AF052692.

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1 - 485	1 - 464	1 - 401	1-197	1 - 108	1 - 440	1 - 211	1 - 385
973358	964722	920365	973359	609722	796358	745133	719663
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HOFMG21	HOFMH12	НОҒМН38	HOFMH95	HOFMI01	HOFMI62	HOFMI63	HOFMJ44

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869113	615305	705435	916588	973351	920218	715312	660317	859104	677372		683473	794308	935553	751692	827631	924473	606999						739399	947431
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HOFNM85	HOFNT59	HOFNU72	HOFNW10	HOFNW79	HOFNY02	HOFNY50	HOFNZ15	HOFNZ16	HOFNZ21		HOFNZ58	HOFNZ94	HOFOA17	HOFOB88	HOFOB91	HOFOE94	HOFOF57				_		HOFOF84	HOGAF39

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1					15 - 629	15 - 1118	15 - 554	15 - 512	15 - 361	15 - 675	15 - 842				15 - 848	15 - 574	15 - 687	
					1 - 615	1 - 1104	1 - 540	1 - 498	1 - 347	1 - 200	1 - 828				1 - 834	1 - 560	1 - 673	
				·	922464	929274	914437	957791	967704	951634	928647				922193	226896	933443	
			·		639	640	641	642	643	645	646				647	648	649	
			,		НР DРQ40	HPDQ005	HPDRB01	HPDRD28	HPDRG92	HPDVB07	HPDVE05				HPDVF03	HPDVK12	HPDVM06	

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AL119522, AL134538, AL119439, AL119484, AL119391, AL119496, AL042970, AL119396,	Ground, microscop, and Albour 1777.		AA299493, AI580853, AA303099, AI287555, AA299492, AI279700, AA559290, N71724, AF177861, AW338869, AA493503, AA367546, AL031777, AC006130, U73024, AC009509, Y10196, U63721, AC006333, AC004047, AJ012197, AL139054, AL050333, AC006544, AC004577, and DR7008	N69328.	AA196263.	T72867, AA610577, and AC007695.	AF111167.			AA737020, AW134485, AI874258. AI217712. AA935591 and AW389859	AC004111.	AI798807, AC007065, AF073930, and AF064804.		AA617957 AT168322 AC005886 AC006071 AC008561 AT008701 AC004332 AC0085730	4 7002004 A 7005156 A 7007300 A 700401 A 700401 A 700401 A 7005010	AC006255. AC004685. AL132987. 293017. AC007308, AL00620114, AP000134, AP000039,	AC004890 AI 040766 A P000555 723840 A COOSESS AI ACOSESS A COOSESS A COOSES A COOSES A COOSESS A COOSESS A COOSESS A COOSESS A COOSESS A	1100001000, ALOUD 1900010, AL VOOLLO, LOSDONO, ALOUD 1009, ALUND 100, ACCOUNTING (CONTRACT) AT 1111115 A COURT OF THE PROPERTY AND ALL OF THE PROPERTY AT 1111115 A COURT OF THE PROPERTY AT 11111115 A COURT OF THE PROPERTY AT 11111115 A COURT OF THE PROPERTY AT 1111115 A COURT OF THE PROPERTY AT 11111115 A COURT OF THE PROPERTY AT 1111115 A COURT OF THE PROPERTY AT 11111115 A COURT OF THE PROPERTY AT 1111115 A COURT OF THE PROPERTY A	230341, ALUSTUSTUSTUSTUSTUSTUSTUSTUSTUSTUSTUSTUSTU	AC008372, AL031587, and AL035249.			AI749823, R99144, AA302658, AW161459, F24175, AI243793, AW161879, AI191343	AA130647, A1679221, A1003797, AA936548, W45457, H24953, AC005057, U62293, U63721	AC005829, AC005102, Z97630, AC005229, AL049830, AF111168, AL035658, AL132712	Z97054, AP000116, AC006241, AC000159, AC002300, AC005747, AB023051, AC009946	AC005323, AC006121, AL049692, AC003030, AC007536, AF047825, AL049757, AC004253.	AC004520, AL109952, AC004030, AL023882, AP000512, U91321, AL022313, AF196779,	AC020663, M55987, AC004893, Z75407, AF102137, AC003665, AC004033, AC002477,	AF20/550, AC012331, AC007666, AF064861, AL049548, Z84480, AC002107, AL030996, AL024507 AC006057 AC004841 AT078644 AC005482 AC	1 ALOCASO1. ACOUNTS A CONTROL OF THE ALTERNATION ACTION AS A CALLA A CALLAND
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	1 - 432	1 - 291	1 - 242	1 - 423	1 - 538	1 - 456	1 - 84	1 - 214	1 - 428	1 - 355	1 - 429	1 - 275	1 - 351	1 - 343							1 - 398	1 - 339	1 - 329								
	955902	867892	208096	723298	655760	716911	914115	922391	921663	969251	960801	524720	965559	236666							655744	009896	781490								
	650	651	652	653	654	655	959	657	658	629	099	199	662	693							664	999	999								
	HPDVY17	HPEAF19	HPEBA06	HPEBA51	HPEBD31	HPEBE44	HPEKG18	HPEK142	HPEKU27	HPEKX12	HPFAA06	HPFCA36	HPFCA71	HPFCF09							HPFCF24	HPFCF40	HPFCF83								

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AC005332, AF165926, AC006061, AC005529, U07561, AC008044, Z95114, X14448, AC004522, AL050307, U63834, AC005846, AL035086, AL022322, AC007226, AP000065, AC005755, AC065914, AC002115, AC005015, AC002073, AC005971, AL031311, AC002350, AC004876, AL022163, Z93930, AC005920, AL031295, AC005527, AC016831, AC005105, AC005081, AC004890, AC002351, AL049780, AC004475, AL035405, AC006449, AC006251, AC008012, AL034379, AC007055, AF001549, AC004475, AL035405, AC003682, AC006512, AC005081, AC005081, AC005081, AC005081, AC005880, AL035450, AC005880, AL031289, AL031289, AL031253, AL031533, AL031659,	AF045555, Z84469, AL022336, AF148808, AC005722, AL049869, AC006948, AC003041, AL022316, AC005184, AC007304, AC005874, AF134471, AP000502, AP000010, Z86090, AC005182, AC005479, AL031685, AL031393, U78027, AL021368, AC007207, AC004815, AC005363, AC002470, Z93017, AL021546, AP000553, Z99716, M63543, AC007917, Z95331, AL121754, AC002395, AC004884, AC005625, AC004889, L78810, AC007021, L44140,	AC005399, AL021878, AL034553, AP000691, AL031230, AC006453, AC002316, AC004024, AC006211, AP000547, AC006515, AL049569, AC002106, and AL031591.				AA651633.	W85782, AA703530, T99014, and C14044.		AI821958, AW150896, and AW205057.				AI525556, AI546855, Z28355, AI526194, AI546999, AI525328, AI556967, AI525316,	A15415/4, 250151, AA585556, C16500, A1525306, A1541508, AA585101, A1541523, AA585439, A1546945, A1546899, A1557799, A1541510, A1525431, A1557731, A1557082	AI541013, AI546875, AI557807, AI546828, AI541534, AI540967, AI526180, AI541307,	AL134524, D61254, AI547039, AI541535, AI526140, AI541365, AI526184, AI536138,	A1557262, T11028, A1540920, A1535660, A1557238, R29445, A1525653, D57491, A1525321,	AL041346, AA585476, AL041096, AL047012, AL041358, AL041377, AL041167, AL041088	AL040621, AL043538, AL041324, AL040464, AL044162, AL041086, AL043496, AL041296,
			15 - 358	15 - 240	15 - 3/0	15-377	15 - 277	15 - 307	15 - 401	15 - 466	15 - 403	15 - 325	15 - 648						
			1 - 344	1 - 226	1 - 356	1 - 363	1 - 263	1 - 293	1 - 387	1 - 452	1 - 389	1-311	1 - 634						
			655768	655538	925750	960365	867871	867880	655749	655621	655533	655563	974257						
			299	899	620	671	672	673	674	675	9/9	229	829				-		
			HPFCH15	HPFCH89	HPFCM87	HPFCN08	HPFCO02	HPFCO67	HPFCQ88	HPFCR21	HPFCR82	HPFCT09	HPFCT53						

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	15 - 324	15-316	15 - 550	15 - 339	15 - 305	15 - 357	15 - 695	15 - 590
·	1 - 310	1 - 302	1 - 536	1 - 325	1 - 291	1 - 343	1 - 681	1 - 576
	655432	655706	525554	655588	960700	655704	974249	973732
	679	089	681	682	683	684	685	989
	HPFCT62	HPFCV19	HPFCV71	HPFCX18	HPFDD06	HPFDE38	HPFDE61	HPFDF79

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		U73330.	AC004111.		AA593648.		AL079301, and AC004530.		AC005225, AL049835, and AL031297.	AW303098, AI061313, AI251203, AI251284, AI250552, AL046519, AI284543, T74524,	AI223626, H07953, AI206841, AI254770, AI189682, AW023111, AI826761, AI357823,	AI251034, AI249853, AA572813, AI375542, AI755202, AI066646, AW410409, AA129746,	AI732430, AA757661, AW276678, AI679759, AW149288, AI635819, AI355007, AW439703,	AI491765, AI753113, AW237905, AI149537, AA702637, AI864500, AI334443, AW192179,	AL038936, AA904211, AIS87583, AA614214, AI247101, AL031730, AC002316, AL031311,	AL022328, AC007055, AL022326, AC006965, AL109865, AF109907, AC005081, AL034343,	AC005592, AL133485, AL049538, AL022238, AC005355, AC005011, AC007358, AC005291,	AL049540, AC008123, AP000688, AC004491, AL080243, AC008372, AL133445, AC003043,	AC004685, AC005874, AF134471, AP000501, AF111167, Z83844, AC004675, AL049780,	AF088219, AC006552, AJ011930, AC007277, AC005330, AC007243, AC006046, AC004686,	AC005363, AC005209, AC005231, AC005746, Z97056, AL135960, AJ131016, AC004883,	AC005619, AC005189, AC002430, AP000511, AL032821, AC006213, AF134726, AC007384,	AC006130, U91326, AC002544, AC004084, AC004067, AL121603, AC005519, AC005899,	AC004821, AF031078, AC008064, AJ010770, AC002369, AF030876, AC009731, AC005844,	AC009510, AL031228, AC005406, AL079352, AC005884, AL022721, AL022476, AL049795,	AC004879, AC005821, AL049760, AC005234, AC004593, AC007129, AC016830, AP000031,	AL023284, X55927, AC006960, AC006203, AC005911, AF001548, AC010202, AC005368,	U07563, AC004542, AL096701, AC006241, Z98750, AC005015, AC004656, AF015416,	AF001549, Z82171, AC002350, AP000134, AP000212, AP000030, U91325, AC007385,	D84401, AC002404, AC007637, AC006511, AC006480, AL031803, AC005606, AC002347,	AC000115, AC005829, AF031076, AL031281, AF053356, AC004051, AL096791, AC005072,	AL049758, AL133353, AC005565, AC016027, AC003101, AC005531, AC004990, AF001552,	AL121652, AC004517, AL117351, AC007011, AC004033, U95742, AC003109, Z70280,	Z83845, AC004147, AC007216, AC004805, AC006079, Z84470, AC005747, AC005225,	ABU23048, ACU03099, ACU03500, ACU04526, ACU05701, D87675, AC004841, AC005722,
15 - 324	15 - 369	15 - 207	15 - 281	15 - 373	15 - 324	15 - 359	15 - 371	15 - 379	15-371	15 - 694			,																						
1-310	1 - 355	1 - 193	1 - 267	1 - 359	1-310	1 - 345	1 - 357	1 - 365	1 - 357	1 - 680																									
655610	655549	954333	655543	581133	655530	974569	522113	867879	739617	867870																									
189	889	689	069	169	692	669	694	695	969	269																									
HPFDG58	HPFDI23	HPFDL90	HPFDS59	HPFDT17	HPFDT54	HPFDT61	HPFDU30	HPFDU38	HPFDU59	HPFDV71																									

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	1 - 641	1 - 328	1 - 386	1 - 295	1 - 213								-									1 - 102					
	655571	655764	960372	925499	953536																	917775					
	869	669	90	701	702																	703					
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	HPMBK49	HPMBM48	HPMBN02	HPMBO10	HPMBO61	HPMBR17	HPMBU81	HPMBX35	HPMBX79	HPMBY76	HPMBY83	HPMBZ05	HPMCB65	HPMCC73	HPMCD77	HPMCI02	HPMCI65	HPMCJ14	HPMCJ19	HPMCJ48	HPMCK65	HPMCS19	HPMCS65	HPMCV93	HPMCW25	HPMCW53	HPMCX11	HPMCY30	HPMCY31	HPMCY35	HPMDJ09	HPMDL78	HPMD039	HPMDR07

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	N69438, N71427, N62439, H80350, R83031, H66851, H67489, R62741, AA918030, H54212, and AC005281	10700011		AC006500,			N20192, and AC008123	AI906494, and AC005004.	AC007966.			AA029876, AA029112, AI089387, W80831, H23039, AA314213, AA209372, AA209368,	1037/02, AASJ4883, AL1189Z/, ALU21/0/, ALU30345, AJ236698, AL080313, and AC006441.	AW 19/446, A19/25/8, A165/152, AW204366, AA436769, AA442937, A1261197, AA421160, AA014360 and A1675734	and the control of th	T58945 T58884 and A1157302	7.07 13, 1200001, alla A12,24.20.	1152.0, A W415101, A W438910, AIZU3120, AAU//81/, AIS65245, F35659, AA318652,	AA221309, N8/410, W60516, AA515938, AW029038, AA025083, A1252554, AA743956,	A1285615, AA618158, AA932099, AW275674, AW236342, A1679045, A1446601, AA177061	F29989, AL133862, AI200051, AI469641, AA631363, AL041368, AA327782, AA626637	H04821, AA745410, A1873916, AL040130, AW440976, AA501600, A1049634 AW008317	AW338500, AA441788, AW069227, AL040913, AI824562. AI633007. AA503454. AA602069	AA228420, AW207652, AI962050, AI358229, AA557879, F36373, AW166815, AW406755	A1985519, A1634187, A1365988, AA713815, AA610786, A1917771, A1457313, A1048035	AI251002, F32705, AI003743, AL042213. AI918421. AW179767. A 8886581. A AA84050	AL040072, AA515051, AW339687, AI246796, A1929531 H65428, A1201124, A1151261	AA515224, AA340960, AI904894, AL120008, AI499094, AW419118, AI635772, A 4 2 2 2 2	AI291268, AI802526, AW069807, AA559290, AI521679, AI537030, AI802023, AI022003	AW162049, AW080134, AI474713, AI433187, AA501722, AI608676, H69041, A A460451	A1049722, U02048, AP000158, S75337, AL035555, Z72521. AF015162, A1031670, AC004057	A39972, U14701, AF140763, AL121603, AF017257, U14684, U14691, U14685, 1114686	U14687, U14688, U14689, U14692, U14699, U14702, U14707, U14710, S75201, U14711	U14712, U14713, AP000014, U14714, U14715, AI 020097, 1114716, AP000692, 1114717
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1 - 334	1-313	1-39	1 - 360	1 - 397	1 - 299	1 - 333	1 - 270	1 - 438	1 - 428	1 - 323	1 - 140	1 - 704	1 201	160-1	1-327	1 - 648	1-220	}											-					
745346	577626	531274	531276	531349	867658	268899	702501	531347	531321	577619	925080	784781	920308	9000	575934	575620	526594		•				_	-										
795	962	797	798	799	800	801	802	803	804	805	908	807	808	 }	809	810	811																	
HPMDT91	HPMDU19	HPMDY82	HPMDZ62	HPMEB66	HPMEC16	HPMEC36	HPMEC69	HPMED52	HPMEE48	HPMEE66	HPMEG50	HPMEI39	HPMFB02		HPMFB28	HPMFB37	HPMFB75																	

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HPMFE35 HPMFF60	813	577633	1 - 415	15 - 429	AC005909,
HPMFE73	815	575941	1-416	15-430	AL009031, and AC002326.
HPMFH21	816	575943	1 - 447	15 - 461	
HPMFJS0 HPMFJS5	818	575932	1 - 475	15 - 489	R20682, T65200, Z44442, and F11983.
HPMFL08	819	929569	1 - 452	15 - 466	AA555286, AA640814, AI281916, AW073979, AI378363, R70468, AW242350, AW013856, AA644290, AW449140, Z93016, AC012384, AL035541, AC005228, AC003662, and AC009300.
HPMFL80	820	874359	1 - 440	15 - 454	AA313895, AA834888, AA362291, AW366931, AA436945, AA496034, AI834310, AA402490, AI039687, and AC004841.

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AA255639, R53909, AI492887, H09775, AW192155, NA88030, 224 A1117508	(1) 1117.000, 1140.000, and ALITON.		AC008072.	AC006121.	R31294, AI823535, AI823533, AL038977, AW271095, A1252141, A1 135383, A1702856	AA594742, AW302832, AA828911, AI284536, AI251563, AI733946, AA578905, AI383596	A1733930, AL047645, A1792695, A1792746, A1792842. A1252190. A1281401 A1689198	T56025, AA553900, AA557493, AA809125, AA484022, AA219181, AI859906, AI089178	AA551067, AI884861, AA169245, AA494047, H81918, AI798313, AI298660, AI038324	A1798569, A1869786, AW269071, A1347649, A1347478, A1761677, AA599080, AC006060	AL022318, AF111168, AL049779, AL034420, X14448, AL021978, AC007731, AC005500,	AJ010598, AC000071, U78027, AC015853, AC005932, AC004033, AF001549, AL035422,	AC006312, AC004263, AC004000, AC005037, AC005412, AL008715, AC005006, AC006057.	AC002544, AC006088, AL133243, AL009051, AL031255, AC005828, Z85996, AC004771.	AC006946, AC004527, AL031577, AC005071, AC004675, AC002470, AC002045, AC002390	AC009263, AL135744, AC004821, AC002300, AC005295, AC005952, AL022316, M26434	AF088219, AL035634, Z98941, Z97053, AC005527, AC005048, AC002106, AC007050	AC008124, AC004262, AC005363, AF015416, AC005730, AC007686, AJ006216, AB017602	AJ246003, AL031133, AF172277, AP000248, AP000704, AL096702, Z81364, AC004912	AL020997, Z83844, AB020868, AL110121, AC006023, AL035681, AL080243, AL008721	AL132987, AL121658, AL049745, AC005368, Z98742, Z85987, AL021391, AC007227,	AF196972, AC000025, AL031428, AL132642, AC005520, AC005041, AL049692, AL031673,	U95742, AL049776, AP000555, AL031846, AL049694, AC004143, AC002115, AC004531,	AC006480, AL033517, AL132777, AC004491, AC002350, AL049872, AC007216, AC004129.	L/8810, AC004797, AL035398, AP000692, AC004941, AC005829, AC005529, L44140,	ALU22119, AL008718, AL020993, US2112, AC007676, AC006064, AC006356, AC002509,	AP000047, AP000226, AF134726, AC004223, AC002477, AC004805, AP000326, Z84466.	AC003689, AF030453, AC007151, AC002394, AC009464, AP000501, Y07950, AC008372.	AC007566, AC004887, AL031602, AC004841, AC004150, AP000054, AP000169, AP000122,	AC004812, AL049569, AC005326, AL109984, AC005323, AL031228, AC004983, AL035416.	AF109907, AP000049, AC005800, AP000115, AL031311, AP000311, AC006241, AC004383,	U85195, AL023876, AP000131, AP000209, AC007435, AP000087, AC005229, AL031433, 798304 AC005088, and 285086	295331.	
15 - 874	15 - 513	15 - 427	15 - 349	15 - 356	15 - 692														<u> </u>	•													15 - 436	15 - 330
1 - 860	1 - 499	1 - 413	1 - 335	1 - 342	1 - 678																				_								1 - 422	1-316
582595	968821	867662	577629	575914	867651																		•							_		-	711640	916662
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HPMFM29	HPMFN12	HPMFP05	HPMFP30	HPMFP38	HPMFQ84																												HPMFS41	HPMFT04

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1 - 326	1 - 330	1 - 363	1 - 353	1 - 266	1 - 466	1 - 436	1 - 341	1 - 346	1 - 348	1 - 512	1 - 308	1 - 367	1 - 421	1 - 422	1 - 796	1 - 551					1 - 355	1 - 402	1 - 387	1 - 269	1 - 347	1 - 414	1 - 492	1 - 517	1 - 648	1 3/2	1 280	1 - 300	1 - 29
575894	652097	575911	956263	960065	932529	575908	867657	577642	577603	968359	506379	920313	703835	867648	873392	954823					577615	577641	924521	920356	864040	575924	671936	730751	854081	\$77500	506235	575051	166676
829	830	831	832	833	834	835	836	837	838	839	840	841	842	843	844	845					846	847	848	849	850	851	852	853	854	855	856	020	/60
HPMFU89	HPMFV28	HPMFV82	HPMFV88	HPMFW25	HPMFW78	HPMFX13	HPMFX65	HPMFX70	HPMFX92	HPMGA83	HPMGB22	HPMGC07	HPMGC23	HPMGE31	HPMGE95	HPMGF06				ocaro, carr	HPMGF32	HPMGH16	HPMGI03	HPMGI84	HPMGJ93	HPMGK37	HPMGK59	HPMGK62	HPMGM33	HPMGM54	HPMGR80	HPMGS00	111 INTOCOL

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				AW242577.		AI806968, AI080168, AI207589, AB006572, and AF091095.	AL133605.	H09672, and A1937462.		AL008709, and AL020990.	AI247199, AW166611, AI254913, AA833896, AA833875, AA644090, AI818737, AI 048466	AW069227, AI460050, R81017, AA613624, D44639, AA603530, AA307598, AW270385	AA634786, AI682665, AA468022, AL047349, AI687343, AI753113, AA528390, AW302017	AA707849, AA984187, AI925579, AA526724, AI084348, AI278972, AI 040054, AI791150	AI867058, AA580808, AA633920, AW188742, AI634187, AI537458, AI004591, AA847409	AA639946, AA574442, AI591375, AA643770, H63092, AW305371, AA651632, AA837035	AA558560, AA862179, AI216990, AW270256, AA846959, AA581247, AI90R093, AA676040	AA084609, AI469577, AI049955, AI720195, AW131043, AA314891, AA714011, AA456974	AA481887, AI732869, N23913, AA191418, AI635279, AW068316, AA452887, H15652	AI283312, AA176605, AA846944, AA904275, AA704393, N64587, AA834777, AA515728.	AA601356, AI457313, AW440545, AI521525, N40092, F30158, AA493226, AA364453	AA583394, AI537020, AA229975, AA521323, AI244127, AA837671, W01985, A1962030	AI580707, AW265138, AA521399, AA086318, AI344948, AA362349, AI302994, AI524360	AA806796, AA721645, AC000025, AC004796, AC005736, AC005081, AC005899, L78810.	AB026906, AC005581, AC004382, AL133448, AC006241, AC004000, U96629, AL096703,	Z98884, AL133245, AL035685, AC006077, AC005844, AC004098, AC007227, AC004876,	AC005746, AC004686, AL078604, AL033517, AL023807, AF196969, 299755, AC005180,	AC005800, AC016025, AL022313, AL132777, AC004253, Z97181, AC004881, Z82172.	AC004973, AC004890, AC002110, AL079305, AC003663, AL132985, AC005632, AC006047.	AC004019, AL031846, AC007327, Z84482, AP000313, AC002104, AC004913, AP000299,	AC005102, AC004447, AC005725, AC000120, AC007666, AP000066, AC006014, Z71183.	AC004476, AP000193, Z99716, U62293, Z93023, AC005619, AC000052, AL034429.	AC005089, AC006441, AL031311, AC005783, AL080243, AC010170, AP000050, AF053356.	AL024507, AF134726, AL035494, AC005768, AL031662, Z83844, AC005231, U47924,	U41193, AC005/40, AF111169, Z81364, AP000117, AP000269, AC005920, AC006197,
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1 - 354	1-1111	1 - 288	1 - 261	1 - 264	1 - 428	1 - 301	1 - 282	1 - 539	1 - 364	1 - 357	1 - 429		•									_	_												
796440	970815	968364	582596	577635	920309	575903	970813	577595	531382	577631	928433				_																				
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HPMGS24	HPMGT67	HPMGV12	HPMGV15	HPMGV59	HPMGW48	HPMGX23	HPMHA80	HPMHB74	HPMHB83	HPMHC74	HPMHD66																								

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							1 - 14/	1-356	1 - 637	1-411	1 - 346	1-310							
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						020	871	872	873	874	875	876							
						TIPN GITTE	HPMJC01	HPMJC05	HPMJD88	HPMJE84	HPMJF76	HPMJ181							

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1	;	15 - 561	15 - 307	15 - 519
		1 - 54/	1 - 293	1 - 505
·	710710	9408/6	922649	917419
·	077	//0	878	879
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	AC006443.	H66737, AA610385, AI825101, and AC006453.		AW130066, AI826186, AW150146, AA129308, AW392049, AA129351, AA453884, AI278397. N24006, AA453799, AI796331, AA336728, N74882, AW380924, AA369050, AI868063,	AW131507, AC004774, D31734, AF033011, U67840, L24443, AF022075, AF022077, AF022076, U25274, AF096161, L09729, U03876, Z63754, and Z63755.	AA457012, and AC002310.	N89001.	AA848128, and Z93244.		R66945, and A1090443.	771183.		A1097067, AI383144, AA370038, and AC005746.	AA370352, AA370860, AA024451, AL118518, AC007016, AF064859, AC004707, Z84718, and	AP000351.	AA370859, AA370351, AC002536, and AL034371.	AA398734, AA393413, and AA371065.	AA371132, AA371091, AW270853, AI978920, AI468384, AI971306, AA361429, and AC005156.	AA371242, AA757085, and AI018367.	and AC002		AC007198.	AA298484.	AA370278.		AC002352.		AA370571, and A1905054.	AA369901,	A1925663, AA458636, AA193435, AA252059, W94787, A1470629, AW016321, and AA193532.
15 - 444	15 - 437	15 - 851	15 - 214	15 - 894		15 - 640	15 - 576	15 - 511	15 - 491	15 - 507	15 - 344	15 - 337	15 - 475	15-329		15 - 228	15 - 466	15 - 242	15 - 332	15 - 300	15 - 352	15 - 320	15 - 298	15 - 335	15 - 357	15 - 279	15 - 272	15 - 419	15 - 306	15 - 581
1 - 430	1 - 423	1 - 837	1 - 200	1 - 880		1 - 626	1 - 562	1 - 497	1 - 477	1 - 493	1 - 330	1 - 323	1 - 461	1-315		1 - 214	1 - 452	1 - 228	1-318	1 - 286	1 - 338	1 - 306	1 - 284	1 - 321	1 - 343	1 - 265	1 - 258	1 - 405	1 - 292	1 - 567
958001	922657	969478	166156	894416		867573	867533	918123	922621	690856	963037	509490	971652	585489		536630	968692	655537	655587	655693	960316	655725	939490	964909	655554	968521	966299	849081	708761	967944
088	881	882	883	884		885	988	887			068	891	892	893		894	895	968	897	868	668	006	901	902	903	904	905	906	907	806
HPMJV08	HPMJY55	HPMKB19	HPMK153	HPMKM81		HPMKN43	HPMLE04	HPMLK02	HPMLK76	HPMLL74	HPMLW10	HPMSF86	HPRAE13	HPRAN84		HPRAU45	HPRAZ10	HPRBA65	HPRBE36	HPRBL91	HPRCB11	HPRCB21	HPRCC08	HPRCC61	HPRCN41	HPRCU13	HPRSB16	HPRTL26	HPRTP73	HPVAB11

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		AIS84019, and R79760.	AA296794.	U18010, A1699466, and L77587.		AW139964, W81697, AW088477, AI887846, AA447817, AA447667, AA677404, R93353.	AA831618, AA574189, AI124782, AA584550, AA055366, F13631, AA765804, AW268271.	AA993918, AA679335, AC008173, AC004074, AF146367, AF124523, AF118808, AL096775,	AL109954, Z83841, AC002464, AC005820, AC006048, AP000509, D84394, AC007359,	AC008175, AC006499, AL136130, AL096801, and W81696.		T65635, A1354862, AA017323, H48816, AI187056, AA829490, AA507745, T47324,	AA548692, H91844, AA565232, AA640022, T08163, N22395, AA524604, AA524829,	AA373304, AI818505, Z83844, AL031255, AC012384, AC004408, AC006449, AC002996,	AC004149, M89651, AC005529, AC004491, AL031228, AC002357, AC002425, AC004531,	AC004947, AC005837, AC004051, AL049779, U52112, AL132774, AC004690, AC005225,	AC006162, AC007686, Z69303, AC006006, AC004231, AC005323, AL136295, AC002511,	AF107045, AC006077, AC004596, AL050317, AL022324, AC006275, AL080245, AL035427,	AL031053, AL049776, AL024507, AC005399, AC004682, AP000688, AC002375, AL078476,	AL121825, AC004552, AC005839, AC007052, AC005527, AL049829, AC007055, AC003049,	AP000555, and AF124730.	AI075673, and AC008015.			T20053.	AW338702.	AW080827, AI535841, AC002038, AC002041, and AC006352.	H68009, AA532955, and AC005189.			AA300733, AW070249, and AC000089.	AW264269, AW028491, AA424983, AA418896, AA418895. AA460211, AI800304, AI685341	AI188340, AI160534, N26011, AW057810, and AI917673.	R59356, F10756, AA480322, AI768068, T15624, AI760446, AI870727, AA534696, AA679825,	Ai86805 1, AI2/4320, AA716748, AI922465, and AA907744.
, 5,	15 - 420	15 - 429	15 - 493	15 - 620	15 - 157	15 - 739				ţ	15 - 227	15 - 344		,								15 - 368	15 - 358	15 - 381	15 - 486	15 - 657	15 - 254	15 - 542	15 - 343	15 - 255	15-410	15 - 799		15 - 152	
717	1-412	1 - 415	1 - 479	1 - 606	1 - 143	1 - 725					1 - 213	1 - 330										1 - 354	1 - 344	1 - 367	1 - 472	1 - 643	1 - 240	1 - 528	1 - 329	1 - 241	1 - 396	1 - 785		1 - 138	
065557	8/5550	753744	525537	655691	655527	908450					655733	867289										657484	655614	095559	655577	514113	961529	537333	655713	165559	710354	705322		785439	
900	20%	910	911	912	913	914					915	916										917	918	919	920	921	922	923	924	925	926	927		928	
UDV/AEA0	III VAL49	HPVAF69	HPVAH36	HPWAF85	HPWAH48	HPWAS77					HPWBA33	HPWBO84										HPWCA53	HPWCJ27	HPWCJ67	HPWCJ82	HPWDA73	HPWDA86	HPWDD72	HPWSB35	HPZAB38	HSWAC73	HSWAD39		HSWBD86	

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AI610767, AA128537, AA045673, H26372, A1564965, C00296, H27484, AA045672, and AF153882.	AI264314, AI656489, AA608739, AL040264, C15165, A1922383, AA644468, AI391663, AI285883, AI538754, AI696671, AA578673, AI587042, AA843485, R27828, AI927062, AA888452, AI379567, AI093793, AA502801, H95376, AI696130, AA032171, AI697448, AA032170, N66388, AI948739, AI499520, and D60629.	AA382253, AI806382, and AI478870.	AA382333, AA382331, H52860, and AL040167.	AA382382, AA382236, and AC006305.	AA382376, AA382242, AA860376, and AC006445.	AA382453.	H53354, AA758164, AA448168, AI025042, AI829657, AA382854, AA382293, AA400727, AA442764, and AA812451.	AA382498.	H09661, AW196402, AA760720, AA432259, AI539284, AI805310, AA781214, AW268890,	AI797250, AW235912, AA993321, AI017538, AW274549, AA884002, AI017535, AI217337,	AW014623, AA868679, AI808721, AI769073, N62418, AI364253, AA909554, AA382799,	A1917688, AA485055, AL080136, and AF079363.	AA429726, AA398189, H53400, AA382855, AA382294, AA383397, AA382295, AA382880,	and Z58143.	AA411883, AW444833, AA382955, AA889519, AI217183, and AA781849.	AA382996, AA180840, AI929073, AL133353, AF132966, and X66366.	AA770016, AW195558, AA757200, AW340425, AW182241, AA383009, and Al191751.	AA383288, AL044302, AA398247, AI916926, AI970265, AA399313, AI806241, AI208574, A 4 904365, A 4 327200, and AI 137385	AA383257, AA812455, and AL137538.	AA383466.	AA383417, and AA383322.	AA383323, AI950478, and AL121757.	AA383746, AF012361, AI918814, AI638690, and AF088868.	AL040588, and AW386895.		AA102609, AA903522, AA810166, AI887877, AA928872, AI912645, AI628759, AI652056,	AA372813, AA326308, AA313595, AA252658, AA491058, AA205648, AW375850, AA252657, and AA082983.
15 - 627	15 - 853	15 - 708	15 - 596	15 - 376	15 - 465	15 - 326	15 - 315	15-311	15 - 425				15 - 680		15 - 760	15-251	15 - 597	15 - 447	15-513	15 - 322	15-317	15 - 471	15 - 1017	15 - 591	15 - 340	15 - 606	
1-613	1 - 839	1 - 694	1 - 582	1 - 362	1 - 451	1 - 312	1 - 301	1 - 297	1-411				1 - 666		1 - 746	1 - 237	1 - 583	1 - 433	1 - 499	1 - 308	1 - 303	1 - 457	1 - 1003	1 - 577	1 - 326	1 - 592	
711066	868007	923070	537136	504353	961232	835805	537271	787500	536769				855596		966113	783328	927010	679390	793441	887782	967959	779134	779265	742218	693377	953904	
929	930	931	932	933	934	935	936	937	938				626		940	941	942	943	944	945	946	947	948	949	950	156	
HSWBJ40	HSWBO34	HTEAA54	HTEAB52	HTEAD32	HTEAD95	HTEAF07	HTEAF26	HTEAG50	HTEAK57				HTEAL28		HTEAP91	HTEAR84	HTEAV43	HTEAY67	HTEAZ54	HTEBC74	HTEBD35	HTEBD40	HTEB178	HTEBP39	HTEBS30	HTEBS77	

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AA300558	AL133028, and AB033037.	M78063, AI560292, H50301, AA340759, AA340760, H45375, 1109355, and AF096153	(COLOC) unit (COCCO) (COCCO)	AB033083.		AA725713, AL040570, R48431, and AC020663	AF193806,	AA382976.	AI017997.	R20356, H09898, and R13239		AI479803, R89200, AA382300, AA700729, and AI 035450				AI521186, AI287890, AW006015, AI739342, AI536021, AI915154, D59412, AI382968,	12(122)/2, C14100, and ICOODO.	41 043427 41100004 4 4 500000 5000000	AD942430, A1123824, AA43708, A1028669, A1024321, AW241753, AA400083, AA953011, AA972296, A1674705, and AL042437.	AW375961, H12646, N46186, AI114644, AA133382, AA314626, T92864, AA482645.	AA203566, W20194, AA878228, W80608, AA699466, AA482495, Z25122, AA490161,	70777101, A1072721, 414 AF 113091.	and AL078621.	AA219332, and AW274715.	AL039822.	268165.	T36107, T19204, AF012359, and T36109	AL079683, T69495, AA377984, AA411067, W19966, AI.135463 AA152011 AIDROTED	A1928496, A1969686, AW150086, AW162108, AW245409, AW245596, AW087350, A1660688.	AW025370, AI986318, AA417624, AW073180, R65681, AA609692, AI568566, AF015913, and AF167572.	AI018016, AA927826, AA781742, AI206323, and AI699610.	
15 - 395	15-418	15 - 358	15 - 266	15 - 623	15 - 244	15 - 435	15 - 354	15 - 407	15 -496	15 - 702	15 - 793	15 - 428	15 - 460	15 - 373	15-315	15 - 1007	15, 331	15 557	100 - 01	15 - 423		15 460	2	15-351	15 - 561	15 - 411	15 - 628	15 - 435			15 - 256	
1 - 381	1 - 404	1 - 344	1 - 252	1 - 609	1 - 230	1 - 421	1 - 340	1 - 393	1 - 482	1 - 688	1 - 779	1 - 414	1 - 446	1 - 359	1 - 301	1 - 993	1 - 317	1 553	1 - 100	1 - 409		1 - 446		1 - 337	1 - 547	1 - 397	1 - 614	1 - 421			1 - 242	
854052	508150	960427	526281	923026	711523	967340	958381	507053	764416	911369	947605	708860	870692	921070	971673	963353	508143	844558	0000	790937		796820		508138	959874	508108	973163	518124	_		527207	
952	953	954	955	926	957	928	959	096	1961	962	963	964	965	996	967	896	696	070	?	971		62.6	!	973	974	975	926	977			8/6	
HTEBS80	HTEBX62	HTEBY08	HTEBY15	HTEBY28	HTEBY41	HTEBY61	HTEBZ21	HTECA13	HTECA16	HTECA21	HTECA32	HTECA51	HTECA83	HTECB21	HTECC13	HTECC20	HTECC37	HTECC38		HTECC66		HTECC80		HTECC85	HTECC96	HTECD17	HTECD18	HTECD36			HTECD62	

HTECD75	979	727422	1 - 410	15 - 424	AA770236.
n i eCeuy	980	020494	1 - 363	15-377	AA074745, AA311579, AW369123, AW377808, AW369199, AA305128, AA279742, AA382491, AA457232, AA350374, and AF119664.
HTECE44	981	764830	1 - 349	15 - 363	AI016755, AA626104, AI215071, AI968379, AA506126, AI340264, AI214879, AI653073, AI208052, AW235456, and AA609852.
HTECE45	982	790894	1 - 349	15 - 363	AA702149, AI760177, W87526, AL042666, AA721639, AA721643, and R55626
HTECE69	983	522983	1 - 340	15 - 354	AA209512, and AF084521.
HTECE91	984	522984	1 - 283	15 - 297	
HTEDF13	985	522964	1 - 370	15 - 384	
HTEDF23	986	522966	1 - 319	15 - 333	
HTEDF57	286	964734	1 - 467	15 - 481	AA206531, AA205273, AF134584, AF133655, R57172, AA370673, AL049834, AF144487, and AF144488
HTEDF76	886	522973	1 - 371	15 - 385	
HTEDG16	686	925527	1 - 1246	15 - 1260	AI341284, AA954970, AA971333, AI656306, AI341232, AI962769, AI970130, AI015246
					A1149570, AW236864, A1187784, A1824998, A1699532, AW087767, AA776716, AA758202, H75549, A1282907, and AC003691.
HTEDG34	066	761752	1 - 601	15 - 615	A65985.
HTEDH21	991	522764	1 - 307	15 - 321	AW237166, and AI806759.
HTEDH22	992	522765	1 - 412	_15 - 426	AA884595, AI004070, AI652358, AI933335, and AC006012.
HTEDH54	993	957762	1 - 398	15 - 412	AI142272, AW303480, and AL117355.
HTEDI02	994	921243	1 - 693	15 - 707	AW166305.
HTEDI16	995	932292	1 - 830	15 - 844	AI200759, N35603, and AI806635.
HTEDI82	966	536477	1 - 536	15 - 550	
HTEDJ04	997	519940	1 - 151	15 - 165	
HTEDJ30	866	771404	1 - 576	15 - 590	AA938733, AI830416, H99099, AI768985, AA410878, AA969071, AI220012, AI568973,
HTEDM08	666	960303	1 - 243	15-257	AA383797 and AA010475
HTED031	1000	870711	1 - 279	15 - 293	
HTEDO51	1001	530592	1-336	15 - 350	
HTEDO59	1002	964379	1 - 277	15 - 291	AL050310.
HTEDP15	1003	870675	1 - 309	15 - 323	AC008119.
HTEDP31	1004	870548	1 - 526	15 - 540	AC004674.
HTEDP32	1005	839532	1 - 412	15 - 426	
HTEDP83	1006	536821	1 - 391	15 - 405	AA382559.
НТЕDQ30	1007	530589	1 - 192	15 - 206	AA470110.

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AA424200		AJ003479, AJ003512, AJ003478, A1719178, and AD000219	AA778548.	AA401417, AI654822, AA131789, AA401393, AA040136, AA030057, AA040089, AW002428, AA131796, AI800387, A1708853, A17088	3, 12, 12, 12, 12, 12, 12, 12, 12, 12, 12		T64832, Z45475, and F17119	AL035450.		AI141062, AA779725, AA451707 AA453324 and AA054530	, 12,12,22,15, 12,12,22,15, allu (AA7,24,22).		AA995142, AC005661, and 265242	AL049780, and AC007055		AI243357, AI126440, AI693753, and A A 783013			AL040261, AI525252 and AI025700	100.000	AA383576, T04917, T35202, AI422683, AI694269, AW341450, AW172708, A1752107	AI392973, AI817020, AI675030, AW149563, AI830691, AI288333, AI420397, AI768573	AW300444, AW271819, A1862664, AW165982, A1761400, A1826256, A1368689, A1697857.	A1889625, A1972408, A1357436, A1421517, A1651095, A1283759, A1634398, A1523543,	A1946511, AW196914, A1992087, A1417903, A1827278, A1927043, AW237591, A1831948,	A1831197, A1830712, AW085599, A1375540, A1300150, A1936396, A1082343, A1190058,	A1569/82, AA93/125, A1926819, AA872799, A1831516, AA502373, A1823952, A1262912, and	A1005426		AI125404. AI247364 AI708717 A A 0 10071 A 1015207	3, 125, 25, 15, 25, 10021, A1910307, and 111493.	
15 - 477	15 - 507	15-317	15 - 377	15 - 1182	15 - 732	15 - 166	15 - 566	15 - 359	15 - 142	15 - 859	15 - 328	15 - 241	15 - 551	15 - 615	15 - 367	15 - 386	_15 - 347	15 - 293	15 - 328	15 - 345	15 - 369							15 - 485	15-312	15 - 876	15 - 148	15 - 224
1 - 463	1 - 493	1 - 303	1 - 363	1 - 1168	1 - 718	1 - 152	1 - 552	1 - 345	1 - 128	1 - 845	1 - 314	1 - 227	1 - 537	1 - 601	1 - 353	1 - 372	1 - 333	1 - 279	1 - 314	1 - 331	1 - 355						-	1 - 471	1 - 298	7 - 862	1 - 134	1 - 210
795332	530586	761585	530451	870708	932315	921114	523959	530452	771505	922964	530580	925399	523962	924818	968517	530157	924840	507814	524059	530199	698315							960127	523957	917206	530095	935982
1008	1009	1010	1011	1012	1013	1014	1015	1016	1017	1018	1019	1020	1021	1022	\neg		1025	1026	1027	-	1029				_			1030		\dashv	-	1034
HTEDQ72	HTEDQ83	HTEDR71	HTEDR91	HTEDU45	HTEDU48	HTEDV02	HTEDX22	HTEDX55	HTEDY38	HTEDY54	HTEDY57	HTEEA03	HTEEB18	HTEEB33	HTEEC10	HTEEE65	HTEEF31	HTEEH31	HTEEU23	HTEEU52	HIEEU88							HTEEU92	HTEEV53	HTEEW73	HTEEZ95	HTEFJ53

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15.463	15 - 494	15 - 151	15 - 418	15 - 303	15 - 412	15 - 353	15 - 322	15 - 196	15 - 734	15 - 358	15 - 341	15-315	15 - 274	15 - 1732										15 - 307	15 - 319	15 - 468	15 - 474	15 - 408			1	15 - 278	15 - 464	15 - 217
1 - 449	1 - 480	1 - 137	1 - 404	1 - 289	1 - 398	1 - 339	1 - 308	1 - 182	1 - 720	1 - 344	1 - 327	1 - 301	1 - 260	1 - 1718										1 - 293	1 - 305	1 - 454	1 - 460	1 - 394				1 - 204	1 - 450	1 - 203
771355	959854	529272	770270	62639	507219	711399	836010	920604	787550	528019	961061	842047	656077	964198						•			2000	/15585	503300	96298	528017	614242			077702	00006/	954115	528007
1035	1036	1037	1038	1039	1040	1041	1042	1043	1044	1045	1046	1047	1048	1049									0201		102	1052	1053	1054			1066	COL	1036	1057
HTEFM31	HTEFNIS	HTEF032	HTEFO76	HTEFP14	HTEFP50	HTEFP61	HTEFS60	HTEFU46	HTEFW55	HTEFW56	HTEFW78	HTEFX53	HTEGA13	HTEGA17									117FF C 4 42	IITECA43	HIEGA4/	HIEGDII	HTEGE44	HTEGF78		_	UTECEOS	TITEOFY	HIEGGO/	HTEGG61

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1 - 759	1 - 406	1 - 526	1 - 544	1 - 438	1 - 300	1-336	1 - 313	1 - 723	1-610	1 - 332	1 - 551	1 - 677	1 757	1-55/							-								1 - 520	1 - 319	1 - 424	1 - 682
870240	573880	794350	783829	917214	573849	870707	526704	973071	866596	573882	795264	784926	022212	716766			_			_					<u> </u>				790342	667224	836999	933624
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1-817	1 - 444	1 - 338	1 - 390	1 - 317	1 - 235	1-317	1 - 430	1 - 414	1 - 390	1 - 664	-1 - 429	1 - 379	1 - 219	1 - 439	1 - 425	1 - 354	1 - 260	1 - 324	1 - 345	1 - 362	1 - 377	1 - 356	1 - 454	1 - 439	1 - 286	1 - 502	1 - 440		1 - 200	1 - 477	1 - 442	1 - 265
935984	526687	233960	922559	870629	530749	528099	771432	573866	573859	787521	920625	967443	531505	573853	751866	573830	573841	529280	573813	786378	870083	920610	785652	924832	760552	924826	772643	┪	\dashv	527167	859130	721831
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	1 - 578	1 - 263	1 - 313	1 - 186	1 - 425	1 - 347	1 - 322	1 - 55	1 - 465	1 - 107	1 - 383	1 - 445	1 - 302	1 - 515	1 - 418	1 - 333	1 - 602	1 - 365	1-322	1 - 344	1 - 253	1 - 356	1 - 576	1 - 504		1 - 444	1 - 535	1 - 390	1 - 859	
	958355	573828	699470	573826	520113	712520	958241	772402	967431	530454	789121	953803	523681	520045	760551	753210	839884	920622	653244	573803	941155	523892	765794	928058		573775	779163	870652	922027	
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	HTEIF40	HTEIF68	HTEIG32	HTEIH70	HTEIJ17	HTEIJ41	HTEIJ73	HTEIJ77	HTEIK11	HTEIK70	HTEIK90	HTEIL07	HTEIL48	HTEIL70	HTEIL71	HTEIN68	HTEIN95	HTEIO02	HTEI012	HTEI028	HTEIP88	HTEIP92	HTEIQ74	HTEIR33		HTEIS65	HTEIU75	HTEIU92	HTEIV54	

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	1 - 476	1 - 790	1 - 340	1 - 725		1 - 364	1 - 365	1 - 342			1 - 640	1 - 373	1 - 336	1 - 364	1 - 326	1 - 292	1 - 412	1 - 577	1-410	1 - 704	1 - 432	1 - 505	1 - 460
	784657	829698	573891	836011		864251	_	577783			955242	523764	528015		870644	573774	772989	908360	520049	942476		774243	573742
	1156	1157	1158	1159		1160	1911				1163	1164		1166	1167	 	1169		1171		1173	1174	1175
	HTEIV86	HTEIW27	HTEIW37	HTEIX28	-	HTEIX85	HTEIY52	HTEIY69			HTEIY80	HTEIZ76	HTEJB20	нтелв25	HTEJB81	HTEJC28	HTEJC95	HTEJE15	HTEJE50	HTEJF45	HTEJG24	HTEJJ43	HTEJL21

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	1-616	1 - 339	1 - 438		1 - 304	1 - 325	1-411		1 - 426	1 - 239	1 - 262	1 - 389	1 - 345		1 - 418	1 - 735	1-1112		1 - 523	1 - 732	1 - 412	1 - 253	1 - 567		1 - 416	1 - 686	1 - 235	1-312	1 - 383
	767955	530596	676254		573823	694525	806395		530156	685272	523818	920925	815975		519938	974044	870084		790381	774260	573750	573749	772997		917176	503039	530588	745257	723148
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1-314	1 - 555	1 - 412	1 - 414	1 - 1063		1 - 333	1 - 495	1 - 426	1 - 490		1 - 286	1 - 304	1 - 304	1 407	1 - 49/	1 - 1005	1 - 510	1 - 495	1 - 601	1 - 442	1 - 441	1 - 251	1 - 658		1 - 575	1 - 654	1 - 376	1 - 453	1 - 227	1 - 646
870627	784444	870555	529273	846714		506651	767658	935945	928656		920927	573700	524054	07672	114708	934272	779315	963563	791743	757740	870621	773018	766462		923071	-	966134	794339	870552	923055
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	HTEMS10	HTEMS48	HTEMT06	HTEMT89	HTEMU17	HTEMUS4	HTEMX92	HTEMY30	HTEMZ04	HTENA22				TITE OF THE PERSON	HIENBUS	HTENC22	HTENF08	HTENF95	HTENG66	HTENG93	HTENH86	HTENI58				UTENIO	TITEINIZO	HIENJ/6	HTENK06	HTENK69	HTEN012	

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AA905347, AA371108, T20204, Z43158, and AA151400.	AA496169, and AC009411.	AL035453.	AA393064.		A1823791, and AF069682.	AI381463, AA634395, AA406053, AA405219, AA383176, and AF121781.	H83100, AL044519, and AL137391.	A1971582, H58143, A1499833, A1393537, and AA973074.	AA225153, AA629286, AA225136, AI272649, AA909816, AA070899, AI866377, AA229443,	R67086, AA084212, AW243884, H59093, AA158549, AC002086, AL133304, AC000004,	AL049830, AC006050, AC007270, AL031846, AC006125, D43727, AL031670, AC002470,	AL050318, AC007114, AF001549, AP000329, AL135959, AC003070, AL034420, AL031286,	AC007226, AL049694, AC004143, AL022476, Z93244, Z68284, AP000359, AC005015,	AL034429, AC000394, AC006120, AC003029, AL031589, AC004782, and AL035072.		AA429691, AA429515, AC007114, Z61140, and AC004156.	AA781188, AA460513, AA860910, AA781845, AI027285, AI208471, AA889700, N73782,	H08088, R38703, F03385, A1830535, A1474644, R41444, AA459870, A1023552, and	AA421081.	AI743533, AI424822, AW082413, and AI915340.	AJ223811, X85630, T36006, AA382232, Z21393, AA383107, N78092, T36070, T36050,	221340, T85719, AL043525, AL043526, R82847, AC006208, and AL137671.	A1018671, A1807205, A1468026, A1797263, A1025828, AW194247, A1889876, AA843455,	AI884356, AI198561, AI032059, AI126485, AI889886, AI239452, AA992969, AA780875,	AW303976, AI216470, AA683361, AI569512, AI472962, AI885458, AW055338, AW242149,	A1911290, A1222107, A1538002, AA912612, AW183126, H79395, AA884115, A1149911,	AW189703, AI220396, AI203939, AA936147, AI560168, AI219573, AW243836, AA906293,	A1252658, N26330, N26296, A1886564, AW085495, AA927058, and AL133596.	AA383398.	AI632084, AI221893, AA383392, AA759214, AW137663, AI269516, and AA923222.	AA383437.	A1990671, and A1990110.	
15 - 649	15 - 508	15 - 496	15 - 655	15 - 589	15 - 458	15 - 889	15 - 546	15 - 1093	15 - 734						15 - 619	15 - 820	15 - 535			15 - 719	15 - 1025		15-976						15 - 630	15 - 578	15 - 503	15 - 566	15 - 562
1-635	1 - 494	1 - 482	1 - 641	1 - 575	1 - 444	1 - 875	1 - 532	1 - 1079	1 - 720						1 - 605	1 - 806	1 - 521			1 - 705	1 - 1011		1 - 962						1 - 616	1 - 564	1 - 489	1 - 552	1 - 548
969213	787535	775387	928244	963530	764828	920834	785996	784936	870515						944416	907717	771409			870587	884043	-	917185						870591	773024	787516	812307	918635
1305	1306	1307	1308	1309	1310	1311	1312	1313	1314						1315	1316	1317			1318	1319		1320						1321	1322	1323	1324	1325
HTENO50	HTENP54	HTENP80	HTENQ05	HTENR10	HTENR74	HTENR93	HTENS22	HTENS43	HTENS91						HTENV57	HTENW53	HTENX77			HTENY21	HTENY35		HTENZ16						HTENZ33	HTENZ72	HTEOA90	HTEOD34	HTEOE61

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AL031283.	AF186084, and AL117610.	AI917508, AI810095, and AA973064.		AA298760, AA298759, AA298601, and AL133073.	AW206247, AI400618, AI671128, AW237084, AI811276, AI337166, AI341186, AI360596,	AW235137, AI183637, AI654814, AI654771, AI969744, AI954759, AA417208, AI969746.	AW087480, AA417104, AW182372, AA723770, AW241414, AW003345, AW241474	AW183339, AA432142, and AA757062.		AW004028, AI968030, AW237673, AA432290, AW138422, AA112090, AA428635, AI143780,	AC004533		AW204420, AW058611, AI767863, AI433866, AI671711, AI802010, AI035766, A A 280370	AI808931, AA523871, AI193461, AA827208 AA418269 AI302011 AAS23871, AA523871, AA523871, AA523871, AA523871, AA523871, AA827208 AA418249 AI302011 AAS23871, AA827208 AA418249 AI302011 AASAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	AI637584, AI932794, AI564719, AI677796, AI538716, AI633125, AI862130, AI432157	A1702073, A1476046, W05052, A1074971 AW15178, AW108000 A1505000 A1501107	A1590277 A1610502 AW102785 A182525 A165525 A1652340 A165232 A1625125	41480118 A1270182 A1067240 A16472019, A10480103, A10480103, A1057240, A1067240	AIG13760 AIG70016 A1371706 A1307331, AW132182, AIG44408, AW104/24, AI863321,	A101273, A1013910, A1211790, A1922363, A1890833, A1926/90, A1802342, A1799199,	At030/19, At241819, At280637, A1696612, A1828731, A1873644, AA833760, A1915291,	A445392, A18 74261, A1804983, A1923357, A1670009, A1560099, A1613017, AL036187,	A1572787, A1570807, AW026882, A1536638, A1620003, A1624548, A1537244, AA449768,	AI433023, AI934026, AW104827, AI537303, AI274508, AI273142, AI955906, AL041772.	AI811785, AI796743, AI446248, AA857306, AI863382, AI554427, AI445025, AI274013,	AI520702, AI446003, AI889376, AW051258, AW150453, AI699865, AI925156, AW151136.	AI923370, AW148320, AI619716, AI539808, AI869367, AI689248, AW075413, AI619607,	AI499285, AW029611, AI698391, AI249962, AI680498, AI567814, AI889189, AI554821.	AI539153, AI554218, AI784252, AW170635, AI608936, AI569945, AI635464, AI432040	AIS37273, AI817552, AI564723, AI612852, AI681985, AI432969, AI828367, AI559296	AI702433, AW075667, AI432030, AI366900, AW148408, AI436644, AI633308, AW169527	AW090550, AI569328, AI627909, AI800411, AI612920, AI919534, AI921176, AI280661.	AI499381, AI469112, AI580984, AI934259, AI539687, AI567128, AW193530, AW073270,	AI648502, AI783861, AI610690, AI889306, AW169653, AI648684, AW081036, AI491775.	AI801523, AW054964, AI469532, AI702068, AW148363, AI819976, AW080992. AW152469
15 - 633	15 - 520	15 - 508	15 - 251	15 - 530	15 - 436				15 - 409	15 - 992	15 - 349	15 - 653	15 - 439					1.					-												
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HTEOF31	HTEOF80	HTEOF85	HTEOF91	HTEOI36	HTE0I53				HTEOK02	HTEON29	HTEON67	HTEOU45	HTEOV90																						

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	15-610	15 - 502	15 - 663	15 - 1068	15-603	15 - 624	15 - 594	15 - 540	15 - 528	15-355								-		-			- -							-	<u>~</u>	_
	1 - 596	1 - 488	1 - 649	1 - 1054	1 - 589	1-610	1 - 580	1 - 526	1 - 514	1 - 341	:											•••••										
	918475	870566	969682	958391	872923	772949	782248	767824	883021	933299									_		•											
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	HTEOW02	HTEOW39	HTEOW85	HTEPA08	HTEPA27	HTEPB66	HTEPB84	HTEPC76	HTEPC87	HTEPD06																						

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	1 - 694	1 - 533	1 - 618	1 - 1202	1 - 453								1 - 359	1-619		1 - 816	1 - 444	1 - 307	1 - 327	1 - 458	1 - 732	1 - 963	1 - 596
	932576	812303	915301	956200	918599								774212	939675		870561	806471	952243	836572	917207	947107	785803	698062
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	HTEPE28	HTEPG15	HTEPH01	HTEPJ09	HTEPJ19								HTEPJ79	HTEPK40		HTEPM33	HTEPM52	HTEPN07	HTEPP23	HTEPP29	HTEPP30	HTEPP32	HTEPP92

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15-169	15 - 407	15 - 644	15-718	15 - 503	15 - 1363	15 - 543	15 - 574		15-410			15 - 557	•		15 - 621	15 - 743	_		15 - 591	15-617	15-510	15-510	15-389	15 - 141	15 - 604	15 - 770	15 - 570				15 - 1312	15 - 361	15 - 267	15 - 334
1 - 155	1 - 393	1 - 630	1 - 704	1 - 489	1 - 1349	1 - 529	1 - 560		1 - 396			1 - 543			1 - 607	1 - 729			1 - 577	1 - 603	1 - 496	1 - 496	1 - 375	1 - 127	1 - 590	1 - 756	1 - 556			1000	1298	1 - 347	1 - 253	1 - 320
787499	870509	870637	915134	917406	869028	963433	888470		922941			915198			918579	958354			881004	853971	870525	806495	806504	966141	932301	939641	966486			0077700	924199	530577	5305/9	575019 1
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HTEPR90	HTEPT25	HTEPT75	HTEPU01	HTEPV02	HTEPX32	HTEPZ10	HTEPZ18	1 C C C C C C C C C C C C C C C C C C C	H1EQB03	٠		HTEQD40		0, 40,441	н ГЕОР69	HTEQE87		THEOCE	HIEUGSO	HTEQI54	HTEQJ14	HTEQJ42	HTEQ081	HTEQP45	HTEQQ82	HTEQR15	HTEQR94			HTEOT62	1111100103	HILABIY	HILAB44	H1LAB/3

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AF038406.		AI217166, AI184534, AI150079, AI142754, AI697160, AI811701, AI214713, and AW188915.		AA418228, AF045454, D63648, and E13935.	AC005258.	AL121757.	AL036680, AA421020, AW151136, R66759, AL037558, AL045421, AA830821, AW149876	AI571529, AI355008, AI567582, AW089006, AI471361, AW023338, AW020095, AI 041150	AW167924, AI829990, AI345224, AI079736, AL046463, AI311892, AW162194, AI697724	T99953, AW243637, AI921167, AA291456, AI539771, AI873638. AI473451. AI805688	AI888621, AI828574, AW022682, AW239367, AI540606, AI689420, AI336662, AL038575	AA464646, AW020693, AI890887, AI590423, AI570966, AI307507, AI916419, AI611728	A1470293, A1929108, A1538850, A1610667, AA572758, A1648567, AW088899, A1805638,	A1366549, AI799195, A1866082, A1636719, A1539153, A1620093, A1866608, A1636619.	AL120853, Al340603, Al537677, Al611743, AW083804, Al349598, Al582912, AW172723	AI539800, AI696626, AI349256, AW075207, AI589993, AI866573, AI312152, AI365256.	AA579232, AA807088, AI343037, AI345735, AW085786, AW265004, AW075084, AI310925	AL038564, AI472536, AI312399, AI677797, AW082600, AI349937, AI567944, AI345688.	A1334884, A1307543, A1494201, N71199, A1345251, AW071412, A1307210, A1307708.	N29277, AI312325, AW071395, AL036631, AI538885, AI249946, AI244380, AI340659,	AIS89267, AW071377, AW129230, AI802240, AW161579, AI313320, AI955906, AI340644.	AI805769, AI434242, AI313352, AI335363, AI307503, AI539707, AI334930, AI309443,	A1623682, AL039086, AI307736, AW161402, AI307520, AI623736, AI446124, AW084097.	A1349266, A1349787, A1334452, A1340664, A1310592, A1344938, A1312146, A1866786,	AJ309431, AJ312339, AJ340537, AW30J300, AJ345739, AW161202, AJ345674, AJ345258,	A1538764, A1312143, A1307459, A1349637, AA635382, A1273179, N74355, A1312428,	A1499974, A1310927, AL110306, A1311604, A1307578, AA420722, AW162189, A1436429,	AI349955, AW189933, AW075093, AI312432, AL120300, AA580663, AI349269, AI312357,	AI590943, AI358701, AW021588, AI310945, AL040241, AW088903, AW151714, AL036638.	AI636581, AI583445, AW059713, AI648408, AI312237, AI922901, AI446373, AW263716,	AI251830, AI343059, AI917963, AI573026, AI349933, AW193467, AW268261. AW082623	AI249877, AL047422, AL133741, AA493923, AI345253, AW409775, AL119836, AI345677,	AW167918, AA494167, AW191003, AI633402, AL119791, AW071362, AW021373, AI345608,	N98606, AW191844, AI336513, AI357599, AI433968, AI499581, AA848053, AI554821,	AW264/19, AL046618, A1348895, A1345347, AW269097, A1866465, A1310575, A1589428,
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HTLCX76	1400	167667	1 - 265	15-279	AL045710, and AL096749.
HTLCY27	1401	682208	1 - 288	15-302	W92263, AA736600, A1962876, A1539379, A1672273, AL045130, A1656897, AW001387, and
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HTLCY54	1402	908832	1 - 1050	15 - 1064	AA453366, AI188219, AI638044, AA983750, AI219830, AA453265, and AA912820.
HTLCZ48	1403	572959	1 - 399	15-413	
HTLCZ96	1404	815897	1 - 459	15 - 473	AW139921, AA725842, AI971598, AI651885, AA453466, AL110422, AI027229, AI026797, AA778573, AI279962, AI073425, AI208767, H55405, and AL118498.
HTLDA58	1405	828115	1-259	15-273	AA948538.
HTLDE53	1406	780842	1 - 610	15 - 624	AA400498, AA400590, AA861265, AA398875, A1149809, A1198885, AA435582, AA460749,
HTLDE64	1407	908613	1 - 838	15 - 852	י משבין התיניטרים, לחינים (הייניטרים), מווח להייניטרים, להייניטרים
HTLDE95	1408	616724	1 - 285	15 - 299	AAS66051, AAS52211, AA393552, AA418209, H05646, H06920, R34733, AA293017, Z42496,
					AI743990, and AA280537.
HTLDF33	1409	909254	1 - 616	15 - 630	A1376558, A1208582, A1149687, A1028197, AA629337, A1027966, A1202003, AA694500, A1701718, AW207552, A1336775, A1560530, A1215526, AW207059, and A1015601.
HTLDG55	1410	911645	1 - 196	15-210	
нтгрн65	1411	839795	1 - 901	15-915	N47989, AI675771, AI024189, AI651589, AI621081, AW003588, AW293606, AI026721,
					AA954294, N51195, and AA983886.
HTLD190	1412	835850	1 - 458	15 - 472	AB020719.
HTLD094	1413	915223	1 - 542	15 - 556	AA860341, AA889689, AA730148, AA478113, AI024743, AA730132, AA082366, and
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HTLDP//	1414	920346	1 - 638	15 - 652	AC002128.
HTLDQ25	1415	870057	1 - 1220	15 - 1234	AA399278, AA889597, AA758803, AW117336, AA398195, AA729781, AA889586, and AA922590.
HTLDSS5	1416	891322	1 - 1302	15 - 1316	AI890919, AI018797, AA913452, AI797580, AI809012, AI187382, AA448485, AI554914, AW137847, AI393577, AA382830, AA432050, AA609003, and AC020663.
HTLDT05	1417	909752	1-473	15-487	R59447, R17647, T62170, and A1439348.
HTLDT81	1418	952265	1 - 480	15 - 494	AA398001, AA399683, AW177623, AA809773, and AL137531.
HTLDU05	1419	911649	1 - 589	15 - 603	AA437044, AF113527, AB023062, and AF113520.
HTLDV31	1420	867748	1 - 297	115 - 311	
HTLDX88	1421	791684	1 - 686	15 - 700	
HTLDY85	1422	573746	1 - 481	15-495	
HTLDZ14	1423	573401	1 - 334	15 - 348	
HTLEB14	1424	573464	1 - 402	15-416	
HTLED72	1425	686906	1 - 273	15 - 287	

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HTLEZ32	1455	870261	1-512	15 - 526	AC005899.
HTLFA74	1456	934172	1 - 695	15 - 709	AA398001, AA399683, AW177623, AA809773. and AL137531.
HTLFC20	1457	917128	1 - 1029	15 - 1043	AA356490, R05624, AA262044, and AL138430.
HTLFE01	1458	917033	1 - 591	15 - 605	
HTLFE05	1459	954984	1 - 1305	15 - 1319	AW275845, AI830267, AI572906, AI439086, AA766499, AA723111, AA836614, AA453028,
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HTLFE20	1460	934278	1 - 463	15 - 477	AC006237.
HTLF128	1461	775392	1 - 423	15 - 437	AC003963.
HTLF139	1462	953730	1 - 616	15 - 630	AF053356, and AF174604.
HTLF183	1463	781303	1 - 501	15-515	AL117491, AB007913, AL137693, and AL110281.
HTLFJ39	1464	573462	1 - 329	15 - 343	AI001797, and AI910520.
HTLGD25	1465	\rightarrow	1 - 1477	15 - 1491	AA608970, and AA758832.
HTLGD69	1466	-+	1 - 1010	15 - 1024	AA403179, AA398239, AW339857, and AA952990.
HTLGG36	1467	789656	1 - 561	15 - 575	AW269804, AA781852, AW104592, AI239999, AI698249, and AC005821.
HTLGK55	1468	868309	1 - 470	15 - 484	AW393087, R16949, R75708, H44330, AA186786, AA259086, R55419, AA329264, AA335735,
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207.0		3133	,		K10335, AA076725, W51754, AL137358, and AL050110.
HTLGM02	1469	964878	1 - 752	15 - 766	C02944, C03248, AI807681, AA095819, AI797565, AI333238, AI863457, H53759, AW138808, AW451889 AW138004 AW1907307 AW184032
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HILG162	1471	918606	1 - 533	15 - 547	AL040693.
HTLGW17	1472	958208	1 - 794	15 - 808	
HTLGX90	1473	870528	1 - 1358	15 - 1372	AA399144, AI982647, AA394141, AA234366, AA405404, AA034080, AA644012, AA234434,
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HTLHC14	1474	908428	1 - 752	15 - 766	AI738485, AI738586, AF096834, AC003688, AF073955, and AF073954.
HTLHE72	1475	963471	1 - 494	15 - 508	AL078621, Z63130, and Z63232.

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1508	509453	1 - 292	15 - 306	AA300600, and AA301000
1509	961057	1 - 247	15 - 261	AA300802, AA300767, and AC004559
1510	797726	1 - 296	15-310	AA300687, AA286758, AL139165, and 770227
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1512	530565	1 - 313	15 - 327	242096.
1513	530562	1 - 252	15 - 266	
1514	530563	1 - 302	15-316	
1515	530564	1-311	15 - 325	AL046237, and AL045777.
1516	578085	1 - 163	15 - 177	
1517	869705	1 - 413	15 - 427	AC005833.
1518	925390	1 - 378	15 - 392	
1519	530393	1 - 112	15 - 126	AL049844.
1520	529672	1 - 239	15 - 253	AA505142.
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				AW118839, R24074, AI803011, AI564140, AI797634, AI671022, AA084054, AA031539,
				A10371233, AA040979, H79028, H03830, AA033515, AW296225, N70304, R31733, AW296783, H30077 A A234000 E32331 IE4138 A A236028
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1522	588965	1 - 203	15 207	AAA20214, AI9/13/0, AA233111, and AA629153.

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	974346	750942	950051	825922	934460	925544	923105	924775			
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974066	931004	953479	926752	869636	869635	934089																-	_	869634	931015	922817	930994	839725	869618	869615	012700	113/99	934130 1		974063 1	\exists	948750 1	
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HTTFZ70	HTTHH05	HTTHU43	HTTIG04	HTTIH23	HTTIH80	HTTIL06																		HTTIN23	HTTIU05	HTTIW81	HTTIZ05	HTTJA11	HTTJA47	HTTJH13	HTTTM01	HTTIONS	11117000	0/24LT-1/11	1111708	H111Y08	HITKD44	

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1-411	1 - 378	1 - 584	1 - 464	1 - 504	1 - 487								1 - 562	1 - 470	1 - 416	1 - 214	1 - 366		1 - 294	1 - 335	1 - 296		1 - 309	1 - 258	1 - 212	1 - 371	1-315		1 - 334	1 - 240		1 - 546		
960928	915033	974311	974316	974310	920893								911390	926795	830008	529157	739638		503626	503445	966804		954420	954359	529727	531165	531163		531110	531108		522213		
1581	1582	1583	1584	1585	1586				_				1587	1588	1589	1590	1651		1592	1593	1594		1595	1596	1597	1598	1599		1600	1601		1602		
HTTKF89	HTTKG34	HTTKK06	HTTKL80	HTTKN21	HTTKN30								HTTKP07	HTTKS13	HTTKV17	HUDAM29	HUDBZ78		HUKAA62	HUKAB80	HUKAC72		HUKAM82	HUKAX07	HUKCC86	HUKDH28	HUKDH50		HUKEH36	HUKEH50		HUKEKSS		

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	15 - 354	15 - 182	15 - 366	15 - 303	15 112	15 - 324		•					·							15 - 344			15 - 280	15 - 367	15 - 372	15 - 386	15 - 331
	1 - 340	1 - 168	1 - 352	1 - 289	1 - 98	1 - 310			-											1 - 330			1 - 200	1 - 353	1 - 358	1 - 372	1 - 317
	714187	967604	526819	537530	946931	575299			_								-			503042			202303	933023	928053	868795	914768
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699896	535361	535006	509119	921088	707061	961020	526248	522823	530558	719332	530387	679477	530386	958027	099898	430750	534783	527937	530094	868779	967815						<u> </u>					
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	1638	1639	1640	1641	1642																							
	HUVFC07	HUVFH03	HUVFH32	HUVF103	HUVFK11			•																				

AC005288, AC004526, AP000359, AP000967, AL109628, AC004882, AC005772, AC002549, AC007878, AC004084, Z99128, AC005324, AC002418, AC003982, AC007314, AL117694, U91318, AC005237, AC002312, Z68226, AC007421, AC008072, AC005041, AC004876, AL031258, AC005740, AC004922, AF001549, AC005763, AL035460, AP000356, AC016027, AL080243, Z69709, AL121655, U80017, AJ246003, AC006130, AP000131, AC002126,	2, and AP000696.	, mil And 1000).		AA076812, F12628, R10205, AA331552, AI752258, R86677, and T74459	, C. L. T.			9. and AB011665.			AA412491, AA412659, AA669386, T93130, T91516, and AB021189	AA658406, AA584733, AL038606, AW189113, AI524360, AI589230, AC006512, AC002994	AC003663, AC006312, AP000104, AC005907, AL109956, AP000272, AC007242, AC006050,	AC005632, AC004675, AP000359, AF088219, U91322, AC003051, and AC005952.			AA120795, AJ003560, AA358990, AW373532, AB014585, AL096767, AC004527, and AC006557		AA477501, AA479873, AI003611, AW270619, AI755214, AI754567. AA584489, A1754105	AI783911, AI962030, AA904275, AI620585, AI570943, AA779783, AA262752, AA013168.	AW277253, AW438856, AL079734, AA582554, AA535216, AA019973, A1491765, A1872216.	AW439703, D26361, AP000493, AC007938, AC004019, AC006210, AC003957, U47924.	AF106564, AC005529, AC006077, AL031311, AF038458, AC006312, AC005578, AC000052,	AL031680, AC004150, AF196779, AC005694, AC007666, AC005295, AC004895, AC005368.	Z93023, AC005755, AC006006, AF053356, AC005682, AC005972, AC005306, Z82172,	AC005231, AC005081, AC007435, AC004972, U62293, Z83844, AC004973, Y18000,	AL049766, AL07887, AP000157, AP000503, AL121653, AC004686, AC004999, AC004812, AL049766, AL078887, AP000157, AP000503, AP0000503, AP00000503, AP0000503, AP0000505, AP0000505, AP0000505, AP0000505, AP0000505, AP0000505, AP0000505, AP0000505, AP00005	AI 121603 AC003006 AI 034402 AE111160 AC014204, AD012303, 227753, Ar 000301,
AC005288, AC00452 AC007878, AC00408. U91318, AC005237, A AL031258, AC00574(AL080243, Z69709, A	AC004813, Z85987, AC005962, and AP000696. A1797913, AA699643, AIS83734, and AA010360.			AA076812, F12628, R	R06515, and AC008116.	AW303494, and AB032978.		AI942325, AI672318, AI672319, and AB011665.	AC005683, and AF172277	H98236, and AC005940	AA412491, AA412655	AA658406, AA584733	AC003663, AC006312	AC005632, AC004675	AC002076.	AA551131.	AA120795, AJ003560,	Z98036.	AA477501, AA479873	AI783911, AI962030, .	AW277253, AW43885	AW439703, D26361, 4	AF106564, AC005529,	AL031680, AC004150,	Z93023, AC005755, A	AC005231, AC005081	AL049766, AL078587.	AT 121603 AC002006
1	15 - 786	15 - 537	15-471	15 - 578	15 - 843	15-371	15 - 422	15 - 384	15 - 356	15 - 402	15 - 418	15 - 159	1		15 - 389	15 - 522	15 - 487	15 - 362	15 - 1165									
	1 - 772	1 - 523	1 - 457	1 - 564	1 - 829	1 - 357	1 - 408	1 - 370	1 - 342	1 - 388	1 - 404	1 - 145			1 - 375	1 - 508	1 - 473	1 - 348	1-1151		_							
	868784	958059	922727	918233	870617	868769	933942	691606	868751	963129	908555	269898			968507	952090	924621	868649	969208									
	1643	1644	1645	1646	1647	1648	1649	1650	1651	1652	1653	1654			1655	1656	1657	1658	1659									_
	HUVFK58	HUVFL71	HUVFQ03	HUVFR02	HUVFT28	HUVFT50	HUVFZ06	HUVGZ77	HUVHB35	HUVHB59	НОУНС93	HUVHD88		2 23	HUVHG80	HUVHI07	НОУНО40	HUVHU74	HVCAZ38									

AC002996, AC007461, AC005224, AC005632, AL049780, Z69917, AC005535, U91321, AF109907, AC007536, Z97183, Z86090, AC004408, AL031602, AC004887, AC004253, AL049697, L44140, AC005520, AL023575, AF134726, AL109657, AP000365, AC007546, AL049776, AC0040313, AC004531, AC005264, AB023051, AD000092, U85195, AC006449, AC003663, AC004055, U95090, AC005620, AC005971, AC002126, AC002395, AC006960, AL031121, AC004890, AE000658, AC004707, AC004472, AC002395, AC006960, AL022322, AP000512, Z83826, AC004407, AC004472, AC002395, AC005914, AC005089, AC009516, AC0040944, AC006512, AP000350, AC003501, AC005089, AC003010, AC00639, AC006639, AC006509, A			AA130954, AI826639, AA149522, AI625572, AI452441, AI653591, AI305234, AI433556, AI366037, AI076932, W60508, AI298406, AW249838, AA693895, AA932571, AI248312, AW136383, AI420053, AW250502, AI216563, AA678309, AA426066, AA937788, F31582, AA340275, AA204645, R11867, AI570112, AI692968, R60411, AI000109, AA130824, AA207010, W56198, ĀW382487, AW382451, AW392670, AW372827, AW363220, AW384394, AL119443, Z99396, AL119391, AL119497, U46351, U46349, AL119483, AL119319, U46347, AL119484, AL119363, AL119457, U46341, AL119341, AL119355, AL119355, AL119439, U46341, AL119341, AL119353, AL119522, AL119418, AL119464, AR066494, Ā81671, AR069079, AR060234, AR054110, and AB026436.					AI751508, AW439242, AI858347, AA954665, AW173684, AA639199, AI633169, AW304822, AI828611, AI088904, AI923258, AI635118, AI952782, AI683485, AI635010, AI963627, AI278264, AW129973, AI860650, AI373090, A647700, AI860650, AI86065
	15 - 632	15 - 63	15 - 917	15 - 273	15 - 493	15 - 389	15 - 555	15 - 1455
	1-618	1-49	1 - 903	1 - 259	1 - 479	1 - 375	1 - 541	1 - 1441
	925844	933449	925796	949141	914354	933675	196896	964102
	1660	1991	1662	1663	1664	1665	1666	1667
	HVCBV04	HVCCC81	HVCCK04	HVVAC41	HVVAJ01	HVVAJ06	HVVBG19	HVVBK45

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	15 - 981	15 - 242	15 - 673	15 - 1615				15 - 300	15 - 622											
	1 - 967	1 - 228	1 - 659	1 - 1601	·			1-286	1 - 608											
	933167	917559	913996	957834				925793	957658											
	1668	1669	1670	1671				1672	1673											
	HVVBK72	HVVBV44	HVVC001	HWLHJ68				HVVDT04	HVVBY08											

AW152182, AI431323, AI862139, AW078729, AI002285, AI932794, AW173633, AW198090, AW192652, AL121007, AI679672, AL046052, AI697207, AI538564, AL046931, AW090550, AI560679, AI884318, AW268964, AI670009, AW264019, AW075936, AI819326, AW006302, AW051258, AI818683, AI783792, AI612852, AI38213, AW129230, AI610671, AI630877, AI921464, AW080746, AI312542, AI890838, AI040725, AL119791, AI473536, AA748343, AI912356, AW168788, AI571439, AI802542, AI922561, AI815855, AI471429, AI570807,	AL037582, AL046595, AI866469, AL037602, AI288050, AI635287, AW026882, AW168031, AI684234, AI559619, AI499963, AI564719, AI951868, AI095119, AI669639, AI890223, AW004886, AI589428, AW169604, AI590021, AI432969, AW078710, AI688858, AW239367, AW079572, AI270183, AI687362, AI499285, AI653979, AI620015, AI522052, AI624693, AI470674, AI67796, AL040207, AI619502, AI978703, AA740450, AI923124, AI682971, AI887396, AW087445, AI284084, AW118477, AI564259, AW148363, AI801766, AW075669.	AW074172, AW088628, AW081298, AI469532, AI872423, AI909661, AI640729, AI283760, AI866770, AW190194, AI679550, AI812015, AI923370, W74529, AI690748, AI613038, AW192687, AI648684, AI539771, AI446809, AI870192, AI538850, AI871697, AI580190, AW129659, AI249877, AI432030, AW083778, AI348914, AI452560, AI679098, AA641818, AI499890, AI890833, AI677646, AI815232, AW081343, AW193020, AI679098, AI241923,	AL387606, AW131294, AW167021, AL934183, AW148408, A1799183, A1537940, AA833906, AC005258, AF182293, Z63483, U77594, Y11587, X65873, X60786, L13297, 189947, AL137271, AF177401, L10353, AF061981, AF113694, U00763, AL080159, X52128, A08910, A08909, 148978, AR038854, 130339, 130334, AL137480, AF111851, AF153205, Z82022, AL137459, AL050149, AL050277, AF139986, A77033, A77035, AL137523, A08908, AF031147, Y09972, 146765, A08916, A08913, U35846, AF067790, AF126247, AL049347, AF111112, AR060234, AF031903, AL049430, AF182215, AL137281, 109499, E05822, AL080110, AL117583, AF090886, AL137533, AL050366, S63521, AL137478, U42766.	M92439, A18788, AF000167, A76335, AF183393, E06743, AL137479, X70685, AF151109, AF085809, AR053103, A65341, AL133557, AF026816, AL137463, AL133093, AF001215, AF111849, AL122100, AF090900, AL080148, A68912, AL137539, X98066, 189931, AL049283, A1000937, AL133075, AL133568, AF104032, 149625, X82434, A18777, AF097996, AL117416, A45787, AF180525, AL133665, S77771, 100734, AL133560, 148979, A57389, AL137526, A1137526,	AL110225, AL110221, MZ 7209, U88909, AR0008994, AL110155, AJ003118, AF138248, AL110225, AF004162, AL137550, AL137488, AF114168, AF100931, U90884, X84990, AF017790, AL137529, Y14314, AL080156, AF090901, AF030513, AL050138, AL137476, AL113437, AF067728, AL133558, Y10936, 133392, AF090903, AL096744, U01145, AL117440, AL096751, AL117435, AL133565, AF032666, Z97214, AF081197, AF081195,
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A. M.	AL137429, AF113677, X87582, AF118090, AL117460, AF026124, U96683, AL023657, AL10077, AL122118, AB026128, Y08769, AS8253, AF078844, AL117648, AL117110, AT660170, AL117648, AR117648, AS826128, AS8253, AF078844, AL117648, AL117648, AR1177110, AT660170, AR1177110,	ALL22110, ALU3011/0, AKULS/9/, AFU/0464, AFU90934, AF114170, L19437, U92068, X99257, ALL133640, Z72491, L30117, AF109155, AL117457, AF036268, X57961, AF057300, AF057299, X83508, AF113690, E03348, AF113689, X66862, AF022813, AR000496, AL137560, AL049382, and U39656.	A1685069, A1378607, A1126015, A1129820, AA912338, A1367538, AW192295, AA495868, and A1738723.	AI761350, AF039240, AI352375, AI206857, T25311, AW291067, AI090441, T25310, W39573, and AC004596.	AW188421, AI810011, AW241649, N51726, AA449707, AI275102, N66707, AI299870,	AI888456, AA912441, N25245, AA909431, H13878, AI362098, H17359, AW020771,	AA858024, AA954574, AA012893, H17330, H08372, H08373, N51811, H08249, H40727,	AA172308. AA247943. AA581747. AA628725. AA354903, AA736863, Z41152, AA172308. AA247943. AA581747. AA019451. H13835. AA33636. AA171007. H15753	AA337210, AI160120, AA854342, AA935605, AW086477, AI242421, T49472, AI471813.	A1803581, AI301683, AI287719, A1803133, AI752956, AA996171, AI983147, AA136130,	AA885946, AA906319, AA429360, AA969237, AI005339, AL037854, AA2471/1, AA910010, AA885946, AA906319, AA429360, AA969237, AI005339, AL037854, AW246747, AA808249	AI282196, AI276053, AA737505, AW015432, AI872692, AA166813, AW250179, AA773516,	A1235032, AA0/0020, A1368453, AWZ/5936, AA608627, A1263811, A1985149, AA833795, AA879424, AI886945, AL037427, AW026473, N25721, AW181174, AW087733, A1 017853	AA553414, AI809312, AI186804, AI005449, AI986314, AI673381, AI017892, N48321,	A1919513, A1468411, AA887602, W60356, AA169201, A1445179, AA554323, AA705759,	A10002/1, A10313/9, A1/42846, AA533/69, AA702089, W45427, AA724285, A1391610, A 8274443 A135060, A1850600, A1420348, A450348, A55048, A450348, A450348, A450348, A450348, A450348, A45048, A4504	AA417981, AW337696, AI453528, AA337233, AI095475, AA17784, AL17748, AL177848, AV337696, AW079854,	D82430, AI363934, AI860112, AA847002, AI743438, AI919040, AI469498, AA577316	TOTAL CONTROL OF THE PROPERTY
			15 - 480	15 - 581	15 - 2282		,												•
			1 - 466	1 - 567	1 - 2268														
			933528	912011	952479						-								
			1674	1675	1676														
			HVVBM06	HUVHL82	HUVHI06									•					•

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AA831676 ACO06318 AEDA4670 2.4 AE157407	AW293310, AI761633, AA721086, AA488897, H86783, AW296923, AA648257, AA906315, H84926. and AF091457.	H15460,	AA368455, AA861915, AA367693, A1000735, D31762, and A1 049611	AA235432.	A1003772, AW304257, AI624133, AW149036, W45510, AW328027, AW069137, AA375087, and D84488			AC004954, and AC004084.	AA664604, AA568204, AA570740, AA483606, AI446623, AA315361 AI 042373 A1569510	H91062, AA449669, W03333, AI801505, N55515, AA603413, AI889579, AI431513, R51582	AI300054, AI253376, AI278089, AW236163, AI356440, AA916177, AA757426, AA130647	AI754170, AI536834, AA513846, AW069412, AA326824, AI627614, T73016, AI784595	AA745470, AA084609, AW406659, AI583106, AI635609, AA401963, AC006129, AJ236701	M87914, S42653, AC006251, AC005368, U62317, AL035587, U96629, AF09572, AC005261	AL080242, AC006468, AC005747, AC005520, AC007283, U80017, AC005585, AL021977	AC007193, AC006449, AC005940, AL022336, AL031283, AC006312, AL031602, AC004999	AC002544, AC002400, AP000240, Z98941, Z94802, AL035458, AC004491, AC006276.	Z98051, AL022165, AC005722, AL031311, AJ011930, AC006064, AC003688, AC005484	AC004814, AC006111, AC004757, AC003080, AP000555, AL031848, U62293, AC000353	AC004815, AC004890, AL135744, AC006071, AC005839, U91318, AC002310, AC005531,	AC007510, AC005086, AC004922, AC004797, AJ246003, AC006285, AC006126, AL022311.	AC006088, U91325, AL031228, AC002996, AC004593, AC000354, AC005412, Z82248,	AC002044, Z84466, AC002418, AL121658, Z98742, AC000086, AL049779, AP000065,	296811, AC005971, Z98304, AC004832, AC005057, AC004386, AF023268, AC002472,	U95742, AC002347, AL035659, Z99943, AC005746, AL121653, U50871, AC005702,	AP000156, AC005212, AC004659, AC002316, AP001053, AC007292, AL034555, AP000096,	AC005519, AC007225, Z84480, AC003109, AL031775, AC007216, AC004881, AJ003147.	AF121781, AC002286, AL021453, AL022320, AC007421, AB022537, AC007707, AF064861	S79349, AL079340, AC002366, AP000014, AC006501, AF124523, AC002115, AC002059	U52112, AL031591, AP000696, Z95115, AP000053, AP000121, AC003966, AC006130,	AC006343, AC000026, AC005823, U33956, AC004106, AL109827, AC005280, AL080243,	AP000251, AC004821, U91326, AP000355, AC009247, AC005031, AC006020, AL022316.	AC005480, AL021579, AC005529, AF196972, AC005763, AC007363, AC007314, AC007430
	15 - 398	15 - 445	15 - 1380	15 - 706	15-313	15-277	15-315	15 - 356	15 - 345								ı																
	1 - 384	1 - 431	1 - 1366	1 - 692	1 - 299	1 - 263	1 - 301	1 - 342	1 - 331						•						•												
	930892	950681	945834	922889	868783	967813	954224	521938	226590	•			_	<u> </u>												-							
	1677	1678	1679	1680	1681	1682	1683	1684	1685																				-				1
	HUVGP05	HUVFP71	HUVF101	HUVFE03	HUVDO18	HUVD011	HUVDO07	HUVDM27	HUVDH61		_																		_			_	

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	15 - 702	15 - 147	15-617									•								
	1 - 688	1 - 702	1 - 603										-							
	974232	671479	667943				_						•		-					
,	1686	1688	1689																	
	HUVCU26	HUVBC21	HUVAA46																	

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AC000477, AC004883, AC004841, AC002326, AL096706, AL031432, AC005796, AC000353, AC000378, AF053356, AC004953, AL139054, AL117337, AC005399, AL050348, AC005606, AC005231, AC004019, AC005562, AC005335, AC005332, AF047825, AC002559, AC006468, AL122003, AL031279, AP000115, AC006449, AP000031, AC007057, AL031602, AL035405, AL021154, AP001053, AC007225, AC004998, AC005664, AC009516, AC005881, and AC005184.	AA757367, AA296644, and AC006327.	R19731, H08375, H06407, R54527, A1654477, AA298292, and A1683415.	AA298987, and AC006222.	AA528216, AI347038, AI308941, AA583432, AA412292, AI147693, AA632915, AA836857,	AA018749, AA135185, AA516435, AA916348, AA514541, N62509, AA470014, AI350971,	R42401, H17978, AA299011, A1207239, AA349833, AA494556, F12523, AA290708, and	R68794, T89302, R64326, R78900, AA299021, AW021631, R26617, and AI820562	AA298749, AW078909, AA742815, AA557686, AC004132, AC008372, AL022238, AC006430.	AC005011, AC005899, AC002351, AF001549, AC004821, AL035420, AC006441, M89651,	AC005746, AJ003147, AL031276, AC005969, AL009031, AC002326, U91318, AC004837,	AC000387, AL034423, Z15025, AC003101, AC005037, AC005184, AC004526, AC018633,	AP000215, AL031295, AL008718, and AP000502.	AA502008, AA298860, N59191, AI559794, AI218041, AI805928, N59181, AW188364,	AA470022, H29622, AI216757, AI015520, AI202594, AA458483, AA565254, AA883777,	N89710, AW340697, AI288448, AA588588, AI344556, AI248194, AI469895, AI954555,	R89606, AI149444, AA505427, AW118040, AA886990, N30779, AW206450, AI800674,	AA343608, AI761704, AA514300, AA581058, and AF151848.	AA258009, AA299129, C02926, N85987, AA630672, AI499588, AW023990, AI636730,	AW196064, AA380354, AA588001, AI818231, AI745151, AL045808, AL046156, AW341978,	AA903253, AA515457, AA308716, AA575911, AA532700, A[494539, AI537772, AL046266,	C15060, AA715201, N53062, AA370455, AI524360, AA487621, AI754544, N94065,	AA488746, AI355556, N23504, AA480574, R73754, AW081194, AA531580, AL138182,	AA618452, W47183, AA580808, AA345064, H69765, F00440, AI684097, AI817516,	AA559241, AI908575, AL119724, R43288, AA833896, AA833875, H12383, AA402129,	AA502991, AI306232, R65651, AA806762, AA604840, AW089322, AI174891, AI002744,	AA463590, AA476397, AA719073, AI623423, H78032, AW085718, AC012627, AL020993,	293017, AF052041, U78308, AF095725, AL121655, AL049631, Z99128, AL035415,	AL035652, AC005377, AF111169, AF003529, AL096766, AJ246003, 295152, AL009179,	AC006353, AL117355, AL121653, AC004655, AC005037, AC006059, AL031279, AL031289,
	15 - 435	15 - 487	15 - 436	15 = 481			15 - 425	15 - 322					15 - 443					15 - 493											
	1 - 421	1 - 473	1 - 422	1 - 467			1-411	1 - 308					1 - 429					1 - 479											
	839574	711543	800452	961527			796691	524239					921132					068896		<u> </u>									
	1690	1691	1692	1693			1694	1695					1696					1697											
	HUNAK12	HUNAG41	HUNAF22	HUNAF20			HUNAE95	HUNAE76					HUNAE02					HUNAD10											

1										
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,	15 - 204	15 - 471	15 - 292	15 - 290	15 - 471	15 - 241				
	1 - 190	1 - 457	1 - 278	1 - 276	1 - 457	1 - 227				
·	753817	968754	714264	754186	574525	760581				
	1698	1699	\neg	-+	-	1703				
	HUNAC68	HUNAB76	HUNAB45	HUKFS69	HUKFL89	HUKFL71				

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i .	15 - 496	15 - 450	15 - 227	15 - 460	15 - 347
	1 - 482	1 - 436	1 - 213	1 - 446	1 - 347
·	574523	577349	574381	796014	921504
	1704	1705	1706	1707	1708
	HUKFL52	HUKFK53	HUKFE56	HUKFD95	HUKER62

					TOTAL COLUMN
111 11/11/11	0.00	72,230			AW015422, Al336885, Z69667, and AC004754.
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					AI536138, T18597, D80045, AI535639, D59751, AI557262, C15076, D80164, AI557084
					D59467, R29657, C14331, D80193, A1526078, A1525856, D81026, C14429, A1541365, D59787
					AA585439, C14389, AI525556, AW366296, D80195, D80227, D59502, AI557533, AI525316
					D50979, D81030, D59275, R45895, D80269, AI541205, D59927, D80022, D80188, D58283.
					C15406, D80166, D51799, D59859, D59619, AI525500, D80210, D80391, D80240, D51423,
					D80253, D80043, AI557864, D59610, AA305409, D80212, D80038, D80196, D80219, D57483.
					A1905856, D80366, D80378, D59889, D50995, AA585098, D52291, D80241, D80251, D80024
					Z32887, AW177440, AI541356, AI557731, Z33559, D51022, AI557602, AA305578.
					AW375405, C14407, AW378532, R28735, R29445, D51060, D80522, R28967, D53161
					D57491, T03269, AW178893, C14014, R28892, AA585378, AA585325, C75569 AA514188
					AW179328, R28965, AI557155, AA585101, T11417, R29218, 732822, D80248, D54897
					D51250, AW369651, AI557751, AI541535, AA514186. AI541346, AI535686 D80134
					AW352158, AI540903, D58253, AI557809, AI540974, AW375406, AW178775, AW178775
					AW177501, AW177511, D80133, AA969188, D80258, AW176467, AW360811, D80133, AA969188, D80258, AW176467
					C05695, D61185, AI557082, AA585356, D60765, F13647, D60844, C14077, AW357117
					AI546829, AW377671, AI557408, AI910186, D80132, AW378540, D80302, AI576184
					AW360844, AW360817, AW378534, AI541034, AW179332, AW377672, AW179023
					AW178905, C06015, AI541517, C16294, T48593, AI546875, AI546999, AI557241, DR0439
					AW352171, Z21582, D59373, AW377676, AW178906, AW352170, AW179018, AI541321
					AW179024, AW177731, AI557734, AI557317, D80247, AW178907, AW179019, D51213.
				,	D81111, AI557787, AW177505, AI546971, AW179020, AW360841, AW178909, D59627.
					AW177456, AW179329, DS1103, AIS57727, AW178980, AIS57852, AW177733, AW378528
					AW178908, AW178754, AI541374, AI526194, AW360834, AW352174, AI536070, AI546945.
					D80014, AW179004, AW179012, AF100707, A62298, A62300, A82595, A84916, AR018138
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		-			A94995, AR031365, D88547, 182448, Y12724, AB002449, AR025207, AF006072, U94592.
	_				AR016514, AR060385, A43190, A44171, AR031358, A30438, Z30183, X68127, AR008443.
	_				A82593, AB012117, AR050070, AR062872, AR038669, I50126, I50132, I50128, I50133,
					A85396, AR066482, AR066488, A85477, AR060138, A45456, A26615, AR052274, A86792.
					X93549, D50010, Y09669, A43192, AR025466, 114842, D13509, AR066490, AR066487.
					118367, AB023656, D88507, AR054175, X76012, AR016691, AR016690, U46128, AR008277.
					AR008281, A63261, AR008408, A70867, AR017826, X82834, 179511, U79457, A64136,
					A68321, AR060133, AF135125, AF213384, AF123263, AB033111, AR032065, AR060382,

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X93535, and AR008382	R54146, and W72450	H56528, and W27624	Z21273, and Z21324	R93336, AI364837, and AI871934	AW188041 AA741890 AA641852 AA654170 AA404652 2-1 AA627400				AA299333 AA299334 AA488228 AI761274 W20642 AA426007 1126851 BC0222	AA442825, AA133785, R48022, N78459, AL046919, A[659107, AA922133, W15736 F07974	AA135871, AW242187, R45109, AI355457, AI937442, AI270750, AI869489, T16709, H04785.	F02239, R36219, AA938012, AI540377, T30366, AA491517, AA897754, AA972881,	A4902408, H19244, Z39481, AI627407, AA133786, AI720129, R56889, AI911499, AI334362.	H16569, A1285897, AI362708, AI148478, AI890959, AI961584, AI625445, AA099547,	A1864641, AA947105, AW295635, AW304383, AW183032, AA580304, A1338026, AA725780,	AA701287, AI815089, AA910649, AA059396, H10768, R47916, AI613048, AI827547,	AA488171, AI362313, R60948, AI351348, AI268730, R69373, AA932084, H17326, H75830	D56341, Z43600, and AA099546.	R02387, and AA299632.	R08548, AI360159, AI445049, AA299487, and AA831602	AA299796, and AA299795.	AA166965, AI143571, AI187051, AI633250, AW135626, AI468530, AI916114, AA299797, and AA299708	AA083174 A1344586 AA000R08 and A1075020	N89673, AA877587, N89666, AW275503, AA548632, AI433318, AI800726, AI013531	AW235557, A1033607, A1741530, AA299661, AW003738, A1431440, D59328, AF108205	AF106698, and U67321.	AA299762, AA299761, AA535406, AA063173, AA534010, M86120. AW069510. AA947547	AI281697, AA310158, F17891, AA584201, AI358431, AA552843, AI367975, AI754955.	D51681, AA846952, N27763, AA367986, AA516226, AA866015, AA488746, D52044	AI653905, AA362349, R89904, AC004142, AL031846, AL031255, U78027, AL031577	AC005237, AC007308, AC004019, AC005578, AC000052, AC005971, AC007216, AC005703.	AP000692, U95742, AL109984, AC005288, AF111168, AC005081, AP000115, AC004234,	ALU22313, ALU20997, AP000553, AC006064, AC004876, AC004895, AL121653, AC005523, AC003562, AC00562, AC00562, AC00562, AC00562, AC00562, A
	15 - 494	15-381	15 - 407	15-310	15 - 450	15 - 350	15 - 260	15 - 126	15 - 399										15-351	15 - 381	15 - 329	15 - 650	15 - 404	15 - 160			15 - 330						
	1 - 480	1 - 367	1 - 393	1 - 296	1 - 436	1 - 336	1 - 246	1-112	1 - 385										1 - 337	1 - 367	1-315	1 - 636	1-390	1 - 146			1 - 316				***		
	920815	719465	968333	690933	967742	529718	529723	518537	628540										502711	502911	672076	502912	716927	790487			524257						<u>.</u>
	1711	1712	1713	1714	1715	1716	1717	1718	1719									300,	1/20	1721	1722	1723	1724	1725			1726						
	HUKDY02	HUKDU47	HUKDG10	HUKCP30	HUKC011	HUKCL85	HUKCC25	HUKCC15	HUKAT59									250 4 200 100	HUKAQ/6	HUKA047	HUKAM19	HUKAM18	HUKAL44	HUKAJ91			HUKAJ83						

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HTTJV79	1743	869602	1 - 434	15 - 448	AA825972.
HTTJN26	1744	869612	1 - 891	15-905	AI356567, N90525, AA232991, AI148171, AI022165, AA233101, AA573721, AA448133,
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HTTIX05	1746	931024	1 - 520	15 - 534	AI493098, AW024745, AA921917, W90789, and AI915946.

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1 - 341	1 - 557	1 - 801	1 - 427	1 - 336		1 - 359	1 - 539	1 - 550	1 - 458			1 - 763	1 - 492	1 - 373	1 - 734		1 - 389		1 - 356				1 - 483		1 - 224	1 - 329	1 - 601	1 - 422			1 - 244		1 - 299
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1 - 576	1 - 475	1 - 301	1 - 378				1 - 583	1 - 456	1 - 356	1 - 627	1 - 550	1 - 509	1 - 602		1 - 713	1 - 472					_		-				1 - 043	1 - 414			1 - 381			1 - 443
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HTEMS01	HTEMO58	HTEMN80	HTEMM91			TTTT: ALC:	HIEMISI	HTEMB57	HTELY90	HTELV29	HTELP07	HTELM71	HTELA02		HTEKU62	HTEKI62										UTTIVITY	/IEWEIL	HTEKD77		707 24 14 14 14	HIEJV94	0.0001	H1EJ046	HTEJN12

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	1 - 1006	1-376	1 - 598			1 474	1 - 4/4	1 - 607	1 - 356	1 - 207	1 - 463		1 - 568	1 - 623		1 - 537	1 - 470	1 - 327	1 - 102	1 - 275	1 - 348	1 - 371	1 - 395	1 - 329			1 - 326	1 - 246
	920628	931017	932987			765001	10600/	732630	709420	545137	866069		887616	666920		719280	685383	530196	707717	575476	530203	530200	675071	677513			780161	530201
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HTECC09	1952	629829	1 - 353	15 - 367	AA306529, AA306591, AA826008, AI089960, AA887528, AI333816, AA513786, AA524465.
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					AA100697, AA070617, AA126067, AA682431, W21262, R58065, AI217985, W95562, H64391,
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					AA281586, AA575968, W95435, AI056604, AI350076, C14543, AA878900, AA947125.
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HTECA44	1954	508135	1 - 371	15 441	AWOREOCE
HTECA39	1955	508132	1 - 408	15 - 422	
HTEBP54	1956	728811	1 - 736	15 - 750	DK3237 U17270 04 D26001
HTEB043	1957	715704	1 - 304	15-318	A1978437 A1791335 A1803274 A7308007 A1095107 A1034518 A 4,00000 A 4,00000
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					AA020798, AA848065, AA436498, AI221817, AI768422. and AA015614
HTEBM87	1958	503275	1 - 278	15 - 292	AA383834, AA383833, AW117332, and AW268492
HTEBL53	1959	578544	1 - 355	15 - 369	AA383796.
HTEBJ02	1960	921321	1 - 408	15 - 422	H73473, and AA383702.
нтевн09	1961	870732	1 - 560	15 - 574	AI204211, AW274576, AI285857, AA383617, AI394396, AA442838 AI218526 AI214386
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15 - 427	15 - 463	15 - 442	15 - 312	15 - 541	15 - 375	15 - 1120					15 - 435	15 - 338	15 - 425	15 - 336	15 - 324	15 - 547	15 - 427	15 - 380	15 - 408	15 - 405			15 - 495	15 - 357			
1 - 413	1 - 449	1 - 428	1 - 298	1 - 527	1 - 361	1 - 1106					1 - 421	1 - 324	1 - 411	1 - 322	1 - 310	1 - 533	1 - 413	1 - 366	1 - 394	1 - 391			1 - 481	1 - 343			
960792	679394	503295	667184	921323	503298	732562			-		503533	503546	503623	960469	724751	925522	708291	867537	412991	953051			666302	471236			
1962	1963	1964	1965	1966	1967	1968					1969	1970	1971	1972	1973	1974	1975	1976	1977	1978			1979	1980			
HTEAX06	HTEAV22	HTEAU39	HTEAT17	HTEAS02	HTEAR93	HTEAQ55					HTEAJ96	HTEAH75	HTEAG47	HTEAG08	HTEAB50	HTEAA04	HSWBY36	HSWBT69	HSWBE29	HSWAS65			HSWAS18	HSWAR63			

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		15-419	15 - 625	15 - 350
	·	1 - 405	1-611	1-336
·	·	697856	936026	727294
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R05355.	W87370, A1168586, AA678331, T96677, W87371, and T96792	R34114, and R62645.	AA036882, AW073808, AW006251, AI992106, AI089385, AI095532, AI143143, AI979301,	AI/30306, AW011022, AII4/683, AA046239, AI398152, AI201784, AW235719, AI696115, and AA011669.	AW172969, AA601278, AA469230, AI376239, AI609972, AA847704, AI431434, AI249365.	AA601680, AA829065, AI053934, AL042539, AA857812, AI281818, AW089016, AI620992,	AA169245, AW089322, AI282629, AA480486, AA704101, AI446623, AA609834, AI859438,	A1340151, AA130647, AA622479, AI912401, W60522, AA904211, AA491767, AI783911,	AA563770, AA487142, AA992562, AI961983, AI417469, AW264901, AA828047, AA82867.	AI133083, AA911590, AA484892, AI224583, AA573062, AW408767, AI279417, AI061313,	AA704393, AL041970, AW407632, AL046519, AI445224, AA197019, T47138, AA578832,	AI445934, AI053597, AI054418, AW303872, AI417464, AA587826, AA487645, AI500552,	H38769, AA176605, AI355559, AA701105, AI636734, AI277373, AI613465, AA362349,	AI989408, AA598605, AI355246, AI733856, W63553, AI598060, AI049931, AI243793.	AW341955, AA985201, AA468956, AI762528, AA632765, AI189682, AA683279, F23338.	AW105729, AI598056, AW157616, R91827, AA515631, AI362552, H95769, AI206841,	AW410409, H71678, AC005740, U80017, U95090, AC006312, AC005696, AC005412,	AC002565, AP000044, AP000692, AC005529, AP000113, AL035587, AL079304, AC004819,	AF111169, AP000045, AL121603, AL035249, AL031311, AL031670, AL139054, AC005081,	AL021546, AC005288, AC005071, AC005730, AL021878, AC004492, AP000555, AC004983,	AP000112, AL035086, AC005952, AC006211, AP000330, AL021391, AL031657, AC005620,	AP000557, AB023049, AC002094, AC004491, AC002115, AC005971, AC007792, AC006111,	AC007057, AL024498, AC004765, AC002316, AC007707, AF196969, AC005225, AC005940,	AC007226, AC000353, AL022238, AF053356, AC002350, AD000092, AC005837, AC005839,	AB003151, AC004796, AL096701, AC004596, AC005037, AF165926, AC020663, AP000501,	AC002312, AP000959, AP000140, AC005231, AC005067, AC004991, AR036572, U91328,	AL078581, AP000031, AC005031, U78027, Z98946, AF001550, AL022318, AF045555,	AC005776, AC005480, AL080243, AC005874, AF134471, AF001549, AF134726, AC006241,	AL050307, AC005409, Z84469, AP000212, AP000134, AC005914, AC004953, AC004910,	AP000350, AC006088, AL035457, AL049759, AC007363, AL109627, D84394, AF111168,	AC004531, AL031846, AC007216, AL034420, AC005803, AC004662, AC005049, AC007686,	AC006049, AC005180, AC003007, AL133445, AL034429, AC006130, AL031281, U47924,	AC003625, AC007298, AL031431, AP000553, AL035659, AC007666, AC006965, AL022326,	293930, AL049/95, AC006011, AC005695, AC002126, AC002302, AC005911, AC005324.
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839559	924978	785710	789170		638155				,														_						_	-	-		-	
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AC004019, AF207550, AL035422, AJ003147, AC004185, AL049709, AC005666, Z83840, AC006285, AL031255, AF064861, AC006543, Z82244, AP000228, Z83845, AC006120, AC004854, AC005200, L44140, AC002477, U52112, AP000088, AL079295, AD000812	AL109801, AC006449, AL031284, AC005520, AL022320, AC004167, AC004253, AC005057, AC003029, U95739, AL021154, AL049692, U85195, AC005829, X87344, AP000248,	AL050341, AL132712, AC009516, AF088219, AC005102, AC006360, AC005082, AC007283, AC007308, AL008718, AC000052, AC004817, 783826, 1191376, AL006791, AC006480	AL020997, AC004929, AC005089, AF030453, AL034548, Z95114, AP000558, Z81369,	AC002425, AL034423, AF196779, AC007450, AC004448, AF196971, AC002404, AC006581,	ACU04000, ALU31584, AC006544, AC005800, AE000658, AC005484, AC002470, 293244, AT117337 AC005470, 293244,	1196629 AL022721 AC005320, AL109796, AC002091, 280090, AL022302, U91318,	AC006254, M89651, AC006538, AC003963, AC007277, AC006255, AC005632, AC007277	AC005722, AC006501, AC002551, and AP000513.	AA136312,	T61306.	W31570, and AF136745.	AI681684, AI458401, AI936264, AW055212, AI200740, AI635739, AI973110, AW044010,	R60842, AI690626, AA281860, AI823929, AI142509, AA781250, and 295115.		N76233, and R18040.	A1671077, AA506744, A1859744, AF047825, AL049757, AC005529, AL022721, AC004491.	AC005823, AC005081, AC005527, AL132712, AL022316, AF001549, AC002312, AC002477,	AC005015, U95742, AP000952, AC005488, AL035587, AP000501, AL021155, AC003982,	AC006511, AF111168, AF109907, AC007216, AC000353, AC000003, AC005484, AC005037,	AC008372, AC005837, 293023, AL035450, AF088219, AC004675, AC004253, AL049761,	ACU06241, ACU05911, AL139054, AL031848, AC009516, AC007227, AC005740, AC005694,	239121, ACUO3000, 233114, ACU01393, ACU023136, ACU024/0, ACU02549, ACU06430, ACU05899 785987 ACU02400 ACU024/0, ACU024/0, ACU02590, ACU024/0, AC	293017, AC005940, AC005696, AI246003, AC002310, AC007050, AL080243,	AC004685, AC006254, AC006547, AC006312, AC005924, L78810, AC007225, AC003108	AP000555, Z84466, AP000553, AC002365, U52112, AC004841, AC005412, AC005565,	284469, AC005291, AL109963, AL034420, AL021546, AP000694, AL121603, and AC006121.		NS9310, and W26272.	AW084140, AI660275, AI159785, AA931541, R46761, AI017335, H69563, AI253649,
					1			1	15 - 233	15 - 546	15 - 291	15 - 340		15 - 216	15 - 565	15 - 497											15 - 285	15 - 392	15 - 672
									1 - 219	1 - 532	1 - 277	1 - 326		1 - 202	1 - 551	1 - 483											1 - 271	1 - 378	1 - 658
									707514	720563	676323	535157		575271	932627	468246											693618	829301	848632
									1989	1990	1991	1992		1993	1994	1995										-+	\dashv	1997	1998
									HPWBF35	HPWBE47	HPWAT23	HPWAJ85		HPWAJ39	HPWAI05	HPWAH19											HFWAG31	HPVAH71	HPVAH41

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	773298	526124	961017	753933	728812	667652	320393	780264	925420	973740	766311	764710	719340	967762	526623											
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	HPVAC74	HPVAB16	HPVAB01	HPRTJ65	HPRTI54	HPRTI16	HPRCV66	HPRCT83	HPRCN03	HPRCM12	HPRCL72	HPRCI73	HPRCG46	HPRCD11	HPRCC22											

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															-							494874	559949	975087	715082	699046	887600	670083		710928	625362	765390	726535
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	,							1 - 385	1 - 372	1 - 358	1 - 235	1 - 303
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HPEBLUS	7104	960240	1 - 408	15 - 422	AI021894, AI359246, AI422145, AI800895, AI689642, T89411, and AC007684
HPEBH01	2165	921767	1 - 333	15 - 347	AA019103.
HPEBG89	2166	785942	1 - 505	15 - 519	AA425851, and AA425691.
HPEBG10	2167	867888	1 - 271	15 - 285	AA533740, and AI984425.
HPEBA89	2168	910250	1 - 490	15 - 504	AA758474,
HPEBA61	2169	742251	1 - 74	15 - 88	AC007036,
HPEBA05	2170	928027	1 - 357	15 - 371	AI217338.
HPEAG43	2171	468542	1 - 485	15 - 499	AA634147, AA837087, N63746, AC006197, AL031311, AC006079, AL035555, AC004790.
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HPEAB12	2173	968847	1 - 498	15 - 512	AI655658, AI991597, AI962317, AW204277, and AI963904
HPEAA57	2174	514231	1 - 441	15 - 455	AC008039.
HPEAA40	2175	867899	1 - 332	15 - 346	AA879011.
HPDWR11	2176	965249	1 - 629	15 - 643	AI694423, AA773635, AA443144, AA443143, AI918138, AW206226. AI337165. AI350411
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HPDWN05	2177	928563	1 - 478	15 - 492	AI627858, AI674644, AI521975, AA653609, AA447670, R37708, H18650, and F34959
HPDVM01	2178	913859	1 - 454	-15 - 468	AA905449, AA969031, U62317, and AC005859.
HPDVD11	2179	965276	1 - 293	15 - 307	AI809869, and AA918215.
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HPDRR06	2181	109896	1 - 436	15 - 450	AA806114, AW242125, AI249498, AI560615, AI274667, AI972210, 728533 and AW361342
HPDRN07	2182	951838	1 - 601	15 - 615	AI365618, AA197089, AW023975, AI570067, AW008217, H05066, AI828721, AI620666.
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	HOVEU10	HOVEU06	HOVEO04	HOVDD82		

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	15 - 405	15 - 618	15 - 586	15 - 469	15 - 443	15 - 318	15 - 335	15 - 1034		15 - 329	15 - 407	15 - 605	707 31	13 - 423	15 - 677					15 - 409	15 - 875	15 - 229	15 - 497	15 - 459	15-411	15 - 486
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	961887	772210	932544	734779	772208	924196	786917	959470		702445	465313	858863	757504	13/394	956238				10000	828857	ᅥ		904818	750273	827077	736077
	2209	2210	2211	2212	2213	2214	2215	2216		2217	2218	2219	2220	2221	7777				3	7777	2223	2224	-	7		2228
	HOVDD10	HOVCP77	HOVCOS0	HOVCN57	HOVCM77	HOVCM03	HOVCI89	HOVCI08		HOVCD33	HOVCC57	HOVBQ07	HOVRV60	TOYOUR	HOVBK38				TOX GIVE	HOVBK 24	HOVBI67	HOVB120	HOVAZ89	HOVAZ65	HOVAY88	HOVAY58

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	AL139054.	AA062554, A1869336, AW170609, H13995, AA631622, H13997, H14028, A1569586, AA632137, and AC004143	H57248.	T79639, N70408, N57728, T83855, AA001363, AI078086, T93794, and W03748	H26739, AI668579, AI733528, and AL020995.	A1031837, A1985078, T25928, AC003692, AL121973, AC004019, AL049766, and AL121603	AA828882, and AA829114.	AI252937, N74027, AA836182, AW338228, AI744524, AA910108, AA772555, A1027046.	A1823533, F02523, A1823535, AW102980, AA745302, AI371165, AW002825, AA887470.	AF001549, AP000502, AC004651, AC002996, U91318, AF134726, AF053356, AC000353,	AL023575, AC004820, AP000113, AP000045, AC004985, AL031587, AC002073, AC005288,	AC005821, AC002310, AC004967, AC005071, Z93930, AP000553, AC007041, AC004703.	AC006014, AC004552, AL023801, AC005015, AL035249, AC005081, AC007371, AC007216.	AP000299, AL136295, AL031230, AL034549, AC003689, AL109628, AC004834, AC007358,	298941, AC005620, AC007955, AC007382, AC007191, AC005765, AC006241, AL020997	AL031283, U91326, AL031659, AC005789, AC002544, AC005837, AL122023, AC005874,	AF134471, AC004814, AC003047, AC005823, AC005086, AC004883, AL049843, AL009181,	AL022323, AL008715, Z98200, AC012330, S79836, U11297, AL049832, AL096791,	AL033547, AC002477, AC004125, AL050317, AC004882, AL031846, AC002357, AC002549,	Z94801, AC007325, AL035405, AC007981, L47234, AC005911, AC000075, U80017,	AC007151, AC002400, AC008079, U78027, AC004531, AC004983, AC004217, AJ003147,	AF045555, AC004837, AC004257, AC006101, AC004953, AC004966, AC005363, AL050325,	AC005606, AC006088, AC020663, AC007688, Z81364, AL024507, AC004832, U95742,	AL022476, Z92542, AC004583, AF039907, AP000208, AC006544, AC005225, AP000247,	AC007229, AL109627, AC008132, AF030453, Z99943, AC002553, Z97183, and AL049748.		R06198.	AA180856, AA312397, AA477002, N64587, F03493, AA722297, AA550850, AI141130.	AW246295, X56997, AC005253, AL133448, AC002477, AC003663, AJ003147, AC007308.	AC006064, AL022320, AC007421, AL022323, AC005632, AC006480, AP000704, Z83840.	AC005786, AL049779, Z85986, AC005520, AC005004, AL132712, Z98036, AC005274,	AC007637, AC005089, AB023049, AC005231, AL049776, AC005180, AL133353, AC007676,	AC009516, AC007298, AC006285, AC005940, AC005081, AL031295, AP000356, AC006211,	ALW21/01, U63/21, AC006101, AC004883, AC005048, AC020663, AC000134, AC012627.
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1.373	1 - 276	1 - 234	1 - 361	1 - 535	1 - 438	1 - 453	1 - 151	1 - 413													-	-			1 763	1 - 333	1 - 499	1 - 410						
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2229	2230	2231	2232	2233	2234	2235	2236	2237																	2230	0000	22.39	2240						
·HOVAY42	HOVAY03	HOVAW62	HOVA025	HOVAN51	HOVAJ07	HOVAI05	HOVAF07	HOVAF01	1								,								HOVAC77	TOVACEA	HOVAC34	HOVAC26						

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	1 - 394	1 - 672	1 - 479	1-577	1 - 345	1 - 541		1 - 472	1 - 238	1 - 372	1 - 430	1 - 423	1 - 415	1 - 795					1-235	1 - 542	1	
	578788	162875	961499	925784	925774	925783		965292	928627	917454	928644	922510	917424	969061					933873	522227	: !	
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	HOVAB85	HOVAB61	HOOKF10	HOOKF04	HOOJ1J04	HOOJN04		HOOJK11	НООЛНОЅ	HOOJE02	HOOIL05	HOOIG03	НООНР02	HOOHE67					90ОНООН	HOOAB23		

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				1 - 569	1 - 324		1 - 1105	_											
				932925	968610		936029									_			
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							1 - 545	.1 - 290	1 - 1041	1 - 595
·							723571	747152	859016	973221
							2259	2260	2261	2262
			,				HONAE50	HONAD65	HONAD02	HOGEW23

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	922578				973227					100037	497/00	463874	964761	829076		930813	859077	161991		745130	750308	847191	_		682232	908904	764490	681919	760431	734848	
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1 - 563	1 - 694	1 - 499	1 - 658	1 - 142	1 - 417	1 - 104	1 - 465			1 - 303		1 - 337			1 - 346
772319	756713	789232	533713	753048	815822	953436	724437	888569	788947	760643	760392	859094	859093	713816	859102
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HOGAD77	HOGAC69	HOFNW81	HOFNW69	HOFNW68	HOFNW65	HOFNW45	HOFNU50	HOFNL96	HOFNI90	HOFNI72	HOFNI71	HOFNI58	HOFNI56	HOFNI42	HOFNI37

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859103	662669	964682	917347	835718	774037	613681								-						731801	725684	489858	743184						-						
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	824242	888552	891512	947973	972725			943358	491360	693987	788733				888780	942367	886485									***		
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	926200	952193	966063	859300	918476	963350	842138			00000	909388	926278	795281	859320	783789	948703	590996	974317	974916	161622	974911	_		_	•					<u> </u>			
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	HODJL04	НОДНК07	HODHG11	HODGL88	HODGJ02	HODFY10	HODFX19			HODEVIA	TODEAIL	HODFX04	HODFW95	HODFW40	HODFR85	HODFL37	HODFJ11	HODFI66	HODFH45	HODFG82	HODFF88		•										

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	926999	909812	774156	974337	915167	952194	963474	859364	779245	859375	745810	531075	765863	420051	920961	745966
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	HODFE04	HODFD73	HODFC79	HODEZ45	HODEU01	HODET07	HODES10	HODE035	HODEK82	HODEA14	HODDX64	HODDS89	HODDS74	HODDF37	HODDE02	HODCZ64

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AC007539.		AA604441, AI690219, AI693458, AA508848, AA447094, AA830582, AI703075, AW003969, AW189292, AW014124, AA831149, AI278956, AI922983, AA908927, AI703009, AA455884, AI870286, AI160778, AI873461, AI636871, A 1745433, AP000504, and AE120756	1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1	H62649, AA584581, AA059154, AL043721, AA098922, AA593060, AA443390, AA329321	C18357, C18360, M77895, AA115968, AA365302, AA658235, AW088224, AI921188,	AA601270, N43757, AA634855, AI286356, AL138455, AL043756, AA515224, AW117750,	AI357551, AI307201, AA630352, AA084070, AI921649, AI929531, A1064952, AI376100,	AW162049, F13749, AA665293, N25303, AA655002, AA641103, AL009179, AL096775,	Z70688, AL022324, AL034419, AL022318, AL078581, AC005962, D87717, AL035696,	U95742, AL049829, AC007068, AC005664, AC000052, AC004019, AF190465, AC004712,	AL023494, AL109627, AL035460, AP000512, AC007022, Z82208, AL122020, AC004797,	AC004622, AC006332, AC004477, AL031005, AL008582, AL008721, AC004859, M87923,	AC005663, AB023051, AC007686, S42659, AC007225, AC004933, AC005406, AC002287.	AC005783, Z95114, AL049776, AL035681, AC006138, AC002119, AC005618, AC004027,	AC006596, AL034548, AL049875, AC005358, AC005911, AF111168, AC005725, AC005288.	AC007546, AC006576, AP000513, AC004526, AC004671, AF072099, L27148, AC004695,	Z82188, AP000270, AC000087, AC004972, AC004927, AP000351, AP000131, AP000209,	U66059, AL031848, AC004638, AC000120, AL078477, AL008629, AC016026, AP000248,	Z98036, AL031782, AC006480, AL133448, AB016897, AL117258, AL023807, AP000032,	AC007690, AC004517, AL031296, AC005324, AC005696, AC006211, AP000349, AC007845,	AC005081, AL022163, AC004035, AL034420, AC005362, AP000049, Z82178, AP000134,	AP000212, AL031291, AP000311, AC007283, AC005808, AL023553, AL031591, AP000080,	AL031846, AC006409, AL109628, AC004522, AC007676, AC004890, AC005104, Z85996,	AC002115, AL022311, AC006132, AF165926, Z97205, AC006285, AC005342, and AF196969.	AW015800, AA251013, and AF037222.	AW304683, and AC004152.	AW104050, AA738352, AI953035, AI341184, AA995803, and AA911220.	AI057634, AW182951, AI655334, AA854636, AI218588, AI457916, AA961245, and	AA953084.	AW007399, AA782657, AI673493, AW104963, AI539419, AI970048, AW272491, AI827847,	AI707847, AI201450, AI160580, AI149344, AI719374, AI870582, AW068652, AW131835,	AA022523, AI142042, AA689495, AI394166, AI197831, AI452812, AA536006, AA483525,	N24911, A1220974, AA994188, AA687451, AA745895, AI808412, A1475847, AA687509,
15 - 321	15 - 146	15 - 77	15 - 201	15 - 833			١				,		_												15 - 327	15 - 450	15 - 706	15 - 596		15 - 509			
1 - 307	1 - 132	1 - 63	1 - 187	1 - 819																					1 - 313	1 - 436	1 - 692	1 - 582		1 - 495			
525832	525834	525833	921187	903653																	_				965121	886650	848179	926973		954374			
2615	2616	2617	2618	2619																				00,0	2620	2621	-	2623		2624			
HBNAC28	HBNAC25	HBNAC09	HBNAC02	HBGTT76				,														_			HBGTEII	HBGQS88	HBGNL85	HBGND04		HBGMT82			

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	15 452	15 - 488		15 - 692	1				,											-											
	1 - 438	1 - 474		1 - 678																											
	754392	773043		880276														•			_									_	1
	2625	2626		2627									-			-							•••					_			
	HBGMS69	HBGMO78		HBGMG81		,														-						•			-		

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AI565755, AI342368, W76561, AA780301, AI200880, AW195697, AA451937, W23209, AA364898, AI050849, H39591, AI028147, AW293972, AI474504, AA324713, AI628717, AA961641, N24530, AF071749, AF136279, AF071748, AF132894, AF088886, AR016587, AL137742, AJ007331, and E15813.		R60478, R11939, AI140927, AW166520, AI097082, AI378818, AW070736, AA022937, AI417973, AI278239, AA813598, AW024249, AI143513, R02801, AW024258, AI097158, AI217615, AA128646, AI831140, AA151765, AI348969, AW058275, AI273856, AI537960, AI310767, AA012905, AL048340, AI952433, AI348854, N22406, AI335235, N20984, AA808904, AI860833, R49697, AI673140, AI810317, AW075382, AL047422, AI288076, AI499104, AI539690, AI473554, AL117403, AF115410, U58996, U68233, I92592, AC005902, AR068751, AF100781, AF213396, Y10080, AL117583, AC004227, AC006288, AF013249, AL133081, AF026124, AF181849, AF181850, AI5345, AL080124, AC005221, AR066486, A70386, AF161699, AF082526, AL049300, I80064, D44497, 108608, AF000301, AF137367, AR023871, AL050310, X56530, and AL110222	T71030, AA747777, AA381858, AA668902, AA558060, T90248, AA934680, AA059235, T70174, AA678436, H86264, AC006001, AC002350, AL021546, AC006449, AC006026, AC001102, AC004895, U18671, AC005387, AL132346, AC022517, AC005696, AL021939, AL035072, AC005089, AC005694, AL050333, AC005565, AL121603, AL135744, AC005839, AL008582, AC005060, U95742, AC006277, AC006071, AL022302, AC006241, AL031681, AC005295, AC005004, AC006277, AC006071, AL022302, AC006241, AL031681, AC005295, AC006530, AC004859, AC000353, A1003147, AC004858, AC005310, AC004253, AC007666, AC006530, AC004859, AC00353, A1003147, AC004858, AC005081, AC002126, AF196969, AC004383, AC00529, AC005940, AC004491, AL022322, AC005015, AP000698, AC005829, AC004030, AC005289, AC005288, Z83840, D00591, AC007308, AL034803, AC005031, AL034420, AC005288, Z83840, D00591, AC007308, AL034803, AC005311, AL049757, AC005288, Z83840, D00591, AC005556, AL031005, AC002369, AL079340, AC002347, AL035659, S77605, Z99716, AP0005556, AL031005, AC002316, AL039340, AC007217, AL035659, AC002218, AC002318, AC002318, AL121655, AC005312, AC0052323, AC005527, AC007193, AC005218, AL121655, AC005550, AL035323, AL034350, AC005535, AC004876, Z83820, AC007021, AL036701, AL032323, AL034350, AC005535, AC004876, Z83820, AC007283, AP000115, AL133371, L78810, AC009516, and AC002316.		
15 - 488	15 - 57	15 - 206	15 - 259	15 - 151	
1 - 474	1 - 43	1 - 192	1 - 245	1 - 137	
832888	802090	588263	522424	525837 971466	
2632	2633	2634	2635	2636	1
HBGDA74	HBGBG69	HBGBG67.	HBGBG52	HBGBG38 HBGBE12	

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H51141, and D37934.	AI306162, AI492835, AC002359, and AC002365.	A1017490, A1221329, AA973064, AA813611, AI688144, A1222206, AW182987, AA883877, W31185, AI208514, AI917508, and AI810095.	AA831288.	AI188098, AW304309, AI375434, AI625524, AI016723, AI167974, AI302664, AW001092,	AI246687, AI572643, AI201622, AW151711, AW167729, AI609516, AI687735, AI191064,	A1289182, A1191358, A1024836, AA632308, A1831665, A1084459, A1088322, AA242818,	AA580006, AI888580, AA011305, AI870130, AW168685, AA010915, AW078807, AI372084,	AI217609, AI200991, AW237859, AI263564, AA629134, AI720347, AA862783, AI880912,	AAS64376, A1879913, A1367291, N74110, Z21520, AA028965, A1349340, AA298576, and A1582057.	R82747, R30969, and T87286.	AA235095, W63805, W07740, AA302624, AA293831, AA315199, R53208, AI568567.	AI992241, AI086596, AW151172, AA309877, AI080304, AA401651, AW090277, AA150647.	AA708776, AA142944, AI826693, AA843315, N77987, AW051448, AA527053, N92512	R87425, A1129519, AI031868, AI083531, AI335895, AI039605, AI299184, AI200329.	AI148099, AA788733, AA115544, AA389673, AW337779, AA588191, AI141373, H66602.	AI192219, AA708520, AA079352, AI076910, AA872411, AA860813, AI160688, AI805893,	AA455203, AA769806, AA411491, AA436345, AA493343, A1193844, A1421247, A1167692,	AI074899, AI193867, AA249309, F32557, AA722032, N80680, W38964, F24549, T83037,	AA402479, AA844466, AI141356, N58395, AA454862, AI161139, AW008542, W93969,	W93970, AA336147, R48412, AI378335, T90357, H95941, AA568295, R86694, AA341105,	AA228455, AW352337, W30921, AW337638, N74117, AA303783, AA215973, AW009811,	R53110, AA558040, AI961636, F24945, AA715148, R49774, F35088, AA228454, AA328964,	AA077962, R11503, H87636, A1695590, AA904185, AA398459, A1803601, R86695,	AA115055, AA192729, AA889647, and AF151893.	AA214667, and H78996.	AA926970, F27522, F30071, AA096277, AA627396, F33364, AA318284, AI283651, F27676,	AW015956, AA947143, AA244132, AA327732, AA352731, AA320725, AA089891, AA317435,	AA319168, F30100, AA664996, F29754, F27644, F27760, AI832645, F24555, AIS60279,	AA306681, AI749712, AI289147, AI130748, AA694164, AI023676, AA244131, F26815,	A1039009, F33405, A1309287, AA534561, T74859, F35771, F34894, F35313, W78719, F30920,	AC008372.	AA127594.
15 - 343	15 - 404	15 - 487	15 - 734	15 - 508				1		15 - 389	15 - 404														15 - 350	15 - 178						15 - 534
1 - 329	1 - 390	1 - 473	1 - 720	1 - 494						1 - 375	1 - 390														1 - 336	1 - 164						1 - 520
773930	932817	927520	922493	917981						951898	764851														577959	518847						685374
2638	2639	2640	2641	2642						2643	2644														2645	2646						2647
HBGBB78	HBCPV80	HBCPO75	HBCPK03	HBCJP02					,	HBCJG07	HAUCC58														HAUAWSI	HAUAS89						HAUAQ28

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T63571, AI832218, T63496, AI004915, AI990727, AW381879, AW381818, AW381815, AI254648. AI917020, and AI 1137755	A1142585, A1082139, AW051999, AA833602, AA719280, AA703255, A1870257, A1952791, AA625853, A1125671, AW104742, AA813022, R44512, AA928937, A1826007, A1921451, A127168, AL120554, A1928736, R48258, A1453416, AW104741, A1933170, AA350537, T98866, AA364915, T40772, A1784134, AA053022, AA740797, AA350621, AA340417, A1240679, AA308837, H16246, AA053159, AA907088, N86840, AW401390, A1939391, AW071136, H30533, R37449, R37338, AC005781, and AC005581	AU292236, AW265393, AI123080, AW069769, AL042373, AW069227, AL038936, AI040051, AW438542, AI216990, AL079734, AA402129, AI887235, AI674946, AA501461, AI733856, AI755214, AW303196, AA469327, AA502532, AW301350, AL120343, AI754105, AA515728, AI754567, AW274349, AI821714, AI792133, AI791913, AW265690, AW270258, AW088846, AW071015, AW372905, AW307312, AI354473, AW1035600, AW270058, AW088846,	AI859946, AI821785, AA653139, AW270270, AL040054, AA613627, AI696962, AI357823, AL042856, AW407578, AA602906, AI801482, AI267356, AW270619, T67090, AI753488, AI281881, AA582554, AW021917, AA634837, AA715814, AI267450, AI874341, AA704393, AL045709, T74524, AA644090, AI754037, AA584489, AI969436, AA631507, AI2822533,	A1054333, AW410354, AW277174, AA470344, AA533025, AW117740, AL044858, AA284247, AA595499, AI473943, AI278089, AW302711, AW270768, AA764783, AI889440, AI275982, AI687343, AI081147, AL042756, AI368256, AW190505, AI291037, AW327624, H07953, AI244127, AW263864, AI371249, AA525253, AA613624, AW438539, AI792499, AI07251, AA8333875, AA833896, AA192278, A1144081, A1380617, A1366580, A167046, A1670648	AI291823, AA430137, AI110844, AA634786, AW020992, AI064864, AW021951, AA307598, AW021154, AI917132, AA622801, H73561, AI872216, AL109758, AC002301, AC006077, AC006077, AC006041, Z82182, AC002306, AC007204, AF205588, AF000251, AC005399, AC006262, AP0000121, AP000030, AC007371, AC006262, AP0000122, AP0000304, AC007371, AC006262, AP0000122, AP0000304, AC007371, AC006262, AP0000124, AP0000304, AC007371, AC007371, AC007808, AC007371, AC007808, AC007371, AC007808, AC007371, AC007808, AC007371, AC007808, AC007371, AC007808, AC007	AC005037, AF053356, AL034417, AC005696, AC002119, AC005261, AL032721, AC004257, AC004815, AC0048	AC005901, AC005519, AC004655, AC002492, AL031433, AL035458, AC006317, AC006966, AC004921, AC004895, AC004477, AL022476, AL133448, AC004858, AC005682, AB023049, AC005060, AC000113, AC005179, AF196972, AC005666, AC008372, AC007055, AL135744, AP00066, AC006676, AC007055, AL135744, AP00066	AC002544, AL121658, AC007684, AC007842, AC002425, AF109907, AL034429, AC002544, AL121658, AC007684, AC007842, AC002544, AL121658, AC007684, AC007842, AC005332, AC005251, AL031228, AC007226, AC007686, AL136504, AF031078, AC004972, AL079340, AC006430, AL033527, AC005701, AF015262, AF165142, AL050307, AC005694,
15 - 527	15 - 453	15 - 499						
1 - 513	1 - 439	1 - 485						
678235	958959	955638						
2648	2649	2650						
HAQCF25	НАQСD07	HACMR08						

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	AC004098, AC009247, Z98200, AC004686, AL139054, AC004128, AL021918, AC004099
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	AC004448, AC007073, AL033525, AC005255, AC004890, AC005324, AC005578, AL031005.
	AC007225, AF111168, AC005726, AC005821, AL035587, AC005207, AC004408, AL022327.
	AC005529, U95742, AC006026, AC004236, AC004150, AC005736, AC007052, AF134726.
	AC000353, AC003108, AF088219, AL022320, AF129756, Z82244, AP000215, AC004263.
	AC016831, AC008115, AF146191, AC004659, AL035405, AL021393, AC005899, AC002365.
	AF001549, AC006101, AL021546, AL117344, Z97056, AL133246, AL034548, Y18000.
	AC003992, AC004707, AF017104, AD000092, AC007899, AC012627, AC005905, AC004590
	AP000505, AC005874, AF134471, AC004796, AL031117, AL109627, AL049643, AL049748
	AC005972, AL049873, AP000252, AL133500, AC004913, AL034555, AC006120, D87675
	AL135960, AJ131016, AC007277, AC005378, AC005071, AC003070, AC004841, AC004622.
•	Z97352, AC002350, AC006254, AC006511, Z98742, AL031276, AC003110, D63507.
	AL096701, AC005216, AC004475, AP000349, Z70288, AL031286, AC005409, AC002073.
	Y10196, AC005520, AC005057, AF064861, AL049843, AC005102, AC004812, AC002418.
	AC005295, AC002553, AL110502, AF190465, AL121603, AC000385, AC007546, AL022162.
	AC005730, AC005484, AC004887, AC004000, AL122023, AC005280, AC003043, AC002504.
	AL031295, AC007537, Z68269, AF015148, AC008044, AB026898, AL109628, and A1420248.
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TABLE 4

Code	Description	Tissue	Organ	Cell Line	Disease	Vector
AR022		a Heart	J.B.	Cen Line	Disease	Vector
AR023	a Liver	a Liver		 		
AR024	a_mammary gland	a_mammary gland				
AR025		a_Prostate	 			
AR026	a small intestine	a small intestine	-		 	
AR027	a Stomach	a Stomach			 	
AR028		Blood B cells				
AR029		Blood B cells		 		
	2.000 2 comb don valed	activated				
AR030	Blood B cells resting	Blood B cells				·
		resting				
AR031	Blood T cells activated	Blood T cells		- 		
		activated				
AR032	Blood T cells resting	Blood T cells resting				
AR033	brain	brain				
AR034	breast	breast				
AR035	breast cancer	breast cancer		- - 	· · · · · · · · · · · · · · · · · · ·	
AR036	Cell Line CAOV3	Cell Line CAOV3		 		
AR037	cell line PA-1	cell line PA-1				
AR038	cell line transformed	cell line transformed				
AR039	colon	colon				
AR040	colon (9808co65R)	colon (9808co65R)				
AR041	colon (9809co15)	colon (9809co15)				ļ
AR042	colon cancer	colon cancer				
AR043	colon cancer (9808co64R)					
711(045	colon cancer (9808c004K)	colon cancer		1 1		ŀ
AR044	colon cancer 9809co14	(9808co64R) colon cancer		 		
7111077	colon cancer 9809cg14	9809co14				
AR045	corn clone 5	corn clone 5	'	+		ļ
AR046	corn clone 6					
AR047	corn clone2	corn clone 6				
AR048	com clone3	corn clone2			·	
AR049	Corn Clone4	corn clone3 Corn Clone4				
AR050	Donor II B Cells 24hrs	Donor II B Cells				
7111050	Bollot II B Cells 241115	24hrs		1		
AR051	Donor II B Cells 72hrs	Donor II B Cells				
	Donot It B Cens /Zins	72hrs				
AR052	Donor II B-Cells 24 hrs.	Donor II B-Celis 24		 		ļ. ———
	Bonor in B-ochs 24 mis.	hrs.				
AR053	Donor II B-Cells 72hrs	Donor II B-Cells				
	20.00 11 2 00.00 72.000	72hrs				
AR054	Donor II Resting B Cells	Donor II Resting B		++	 -	
	- and a comp	Cells				
AR055	Heart	Heart		 		
AR056	Human Lung (clonetech)	Human Lung		 		
	Lang (cronetten)	(clonetech)				
AR057	Human Mammary	Human Mammary		+	-	
	(clontech)	(clontech)				
AR058	Human Thymus	Human Thymus		- -		
I	(clonetech)	(clonetech)				
	(CloudleCecii)	(5.5.1510011)				
		Inrket				
	Jurkat (unstimulated)	Jurkat (unstimulated)]]	ł	
AR059	Jurkat (unstimulated)	(unstimulated)				
AR059 AR060	Jurkat (unstimulated) Kidney	(unstimulated) Kidney				
AR059 AR060 AR061	Jurkat (unstimulated) Kidney Liver	(unstimulated) Kidney Liver	<u>-</u>			
AR059 AR060 AR061 AR062 AR063	Jurkat (unstimulated) Kidney	(unstimulated) Kidney				

AR064				-,			
B cell lymphoma	AR064	Lymphocytes diffuse lease	leukaemia	 			
AR065 Jymphocytes follicular Jymphoma	AKUU4	- July 100 dillase laige	1 ' ' '			İ	
AR060 Lymphocytes follicular Lymphocytes Lymphocyt		B cell lymphoma		1 .]
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AR062	AR066			 			
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AR069 Normal Ovary 9701G208 Normal Ovary 9701G208	A D 0 6 8					<u></u>	
AR069 Normal Ovary 9701 G208 Normal Ovary 9701 G208 AR070 Normal Ovary 9806G005 9701 G208 AR071 Ovarian Cancer	AKUU	Normal Ovary 9508G045			1		
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AR070 Normal Ovary 9806G005 Normal Ovary 9806G005	AKOOS	Normal Ovary 9701G208					
AR071	A P.070	No-mal Over 1000 C005		ļ			
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AR095 Small Intestine (Clontech) Small Intestine			•			ł	
	AR095	Small Intestine (Clontech)					

L D006	C-1					
AR096	Spleen	Spleen	ļ			
AR097	Thymus T cells activated	Thymus T cells activated		,		
AR098	Thymus T cells resting	Thymus T cells				
AR099	Tonsil	resting	 			
		Tonsil			ļ	
AR100	Tonsil geminal center centroblast	Tonsil geminal center centroblast				
AR101	Tonsil germinal center B	Tonsil germinal center B cell				
AR102	Tonsil lymph node	Tonsil lymph node	 	 	 	
AR103	Tonsil memory B cell	Tonsil memory B				
AR104	Whole Brain	cell	 -	 -		<u> </u>
AR105	Xenograft ES-2	Whole Brain	 	ļ	_	
AR106	Xenograft SW626	Xenograft ES-2	ļ		<u> </u>	
H0008	Whole 6 Week Old	Xenograft SW626 ,		ļ	ļ	<u> </u>
	Embryo					Uni-ZAP XR
H0009	Human Fetal Brain					Uni-ZAP XR
H0012	Human Fetal Kidney	Human Fetal Kidney	Kidney			Uni-ZAP XR
H0013	Human 8 Week Whole Embryo	Human 8 Week Old Embryo	Embryo			Uni-ZAP XR
H0028	Human Old Ovary	Human Old Ovary	Ovary		† 	pBluescript
H0030	Human Placenta				 	Uni-ZAP XR
H0031	Human Placenta	Human Placenta	Placenta		 	Uni-ZAP XR
H0032	Human Prostate	Human Prostate	Prostate			Uni-ZAP XR
H0038	Human Testes	Human Testes	Testis	 	 	Uni-ZAP XR
H0040	Human Testes Tumor	Human Testes Tumor	Testis		disease	Uni-ZAP XR
H0046	Human Endometrial Tumor	Human Endometrial Tumor	Uterus	1	disease	Uni-ZAP XR
H0050	Human Fetal Heart	Human Fetal Heart	11	ļ ·		
H0051	Human Hippocampus	Human	Heart			Uni-ZAP XR
	Truman Trippocampus	Hippocampus	Brain			Uni-ZAP XR
H0052	Human Cerebellum	Human Cerebellum	Brain			Uni-ZAP XR
H0055	Human Umbilical Vein	Human Umbilical	Umbilical			Uni-ZAP XR
		Vein Endothelial Cells	vein			OIII-ZAI AK
H0056	Human Umbilical Vein,	Human Umbilical	Umbilical		<u> </u>	Uni-ZAP XR
	Endo. remake	Vein Endothelial Cells	vein			OIII-ZAI XX
H0057	Human Fetal Spleen					Uni-ZAP XR
H0059	Human Uterine Cancer	Human Uterine Cancer	Uterus		disease	Lambda ZAP II
H0063	Human Thymus	Human Thymus	Thymus			Uni-ZAP XR
H0087	Human Thymus	Human Thymus	inymus	 -		
H0090	Human T-Cell Lymphoma	T-Cell Lymphoma	T-Cell	 -	disease	pBluescript
H0102	Human Whole 6 Week	Human Whole Six	Embryo	 -	uisease	Uni-ZAP XR
	Old Embryo (II), subt	Week Old Embryo				pBluescript
H0111	Human Placenta, subtracted	Human Placenta	Placenta			pBluescript
H0124	Human Rhabdomyosarcoma	Human Rhabdomyosarcoma	Sk Muscle		disease	Uni-ZAP XR
H0144	Nine Week Old Early Stage Human	9 Wk Old Early Stage Human	Embryo		····	Uni-ZAP XR
H0150	Human Epididymus	Epididymis	Testis			Ilmi ZADVO
H0156	Human Adrenal Gland	Human Adrenal	Adrenal		disease	Uni-ZAP XR
	Tumor	Gland Tumor	Gland		uisease	Uni-ZAP XR
H0163	Human Synovium	Human Synovium	Synovium			Uni-ZAP XR
H0165	Human Prostate Cancer, Stage B2	Human Prostate Cancer, stage B2	Prostate		disease	Uni-ZAP XR

110166	4.1.					
H0166		Human Prostate	Prostate		discase	Uni-ZAP XR
H0168	Stage B2 fraction	Cancer, stage B2				
HOTOS	Transfer Transfer Culture,	Human Prostate	Prostate		disease	Uni-ZAP XR
110160	Stage C	Cancer, stage C				
H0169		Human Prostate	Prostate		disease	Uni-ZAP XR
110170	Stage C fraction	Cancer, stage C				1
H0170	- Tolk old Barry Stage	Twelve Week Old	Embryo			Uni-ZAP XR
11015	Human	Early Stage Human			j	
H0171	12 Week Old Early Stage	Twelve Week Old	Embryo			Uni-ZAP XR
	Human, II	Early Stage Human			į.	,
H0176	CAMA1Ee Cell Line	CAMA1Ee Cell	Breast	Cell Line	1.	Uni-ZAP XR
		Line	<u> </u>		1	
H0178	Human Fetal Brain	Human Fetal Brain	Brain			Uni-ZAP XR
H0179	Human Neutrophil	Human Neutrophil	Blood	Cell Line		Uni-ZAP XR
H0181	Human Primary Breast	Human Primary	Breast		disease	Uni-ZAP XR
	Cancer	Breast Cancer		1		Om Zan Aik
H0182	Human Primary Breast	Human Primary	Breast	†	discase	Uni-ZAP XR
	Cancer	Breast Cancer			, discuse	OIII-ZAI AK
H0188	Human Normal Breast	Human Normal	Breast		i — —	Uni-ZAP XR
		Breast	1	١,		Olli-ZAI AK
H0194	Human Cerebellum,	Human Cerebellum	Brain		†··	pBluescript
Ĺ	subtracted					polacscript
H0196	Human Cardiomyopathy,	Human	Heart			Uni-ZAP XR
<u> </u>	subtracted	Cardiomyopathy	110411	İ		UIII-ZAF AK
H0211	Human	Human Prostate	Prostate	 		pBluescript
Į	Prostate, differential		1105.010		1	poluescript
L	expression			1	1	1
H0212	Human Prostate,	Human Prostate	Prostate		 	- Diversity
	subtracted		1 Tostate	1	i	pBluescript
H0244	Human 8 Week Whole	Human 8 Week Old	Embryo	 		LU-17AD VD
	Embryo, subtracted	Embryo	Linuiyo			Uni-ZAP XR
H0252	Human Osteosarcoma	Human	Bone		disease	11-: 740 20
		Osteosarcoma	Done		disease	Uni-ZAP XR
H0253	Human adult testis, large	Human Adult Testis	Testis			11 : 7 + 7 1/7
	inserts	Transact reduct resting	103013			Uni-ZAP XR
H0255	breast lymph node CDNA	Breast Lymph Node	Lymph Node	 		1 1
ļ	library	Droubt Lymph Houc	Lymph Node]		Lambda
H0263	human colon cancer	Human Colon	Colon		4:	ZAP II
		Cancer	Colon		disease	Lambda
H0266	Human Microvascular	HMEC	Vein	Cell Line		ZAP II
	Endothelial Cells, fract. A	INVILO	A CIL	Cell Line		Lambda
H0270	HPAS (human pancreas,	Human Pancreas	Pancreas			ZAPII
1	subtracted)	Tunian Fancicas	Pancreas			Uni-ZAP XR
H0271	Human Neutrophil,	Human Neutrophil -	Blood	0.11.1		
110271	Activated	Activated	B000	Cell Line		Uni-ZAP XR
H0294	Amniotic Cells - TNF		Di			
110274	induced	Amniotic Cells - TNF induced	Placenta	Cell Line		Uni-ZAP XR
H0295	Amniotic Cells - Primary	Amniotic Cells -		<u> </u>		
110273	Culture		Placenta	Cell Line		Uni-ZAP XR
H0310	human caudate nucleus	Primary Culture	D.:	 		
H0316	HUMAN STOMACH	Brain	Brain			Uni-ZAP XR
H0318		Human Stomach	Stomach			Uni-ZAP XR
H0328	human ovarian cancer	Ovarian Cancer	Ovary		disease	Uni-ZAP XR
TU341	Bone Marrow Cell Line	Bone Marrow Cell	Bone Marrow	Cell Line		Uni-ZAP XR
U02/0	(RS4;11)	Line RS4;11				
H0369	H. Atrophic Endometrium	Atrophic		[Uni-ZAP XR
}		Endometrium and		· }	1	1
770255	11	myometrium				
H0372	Human Testes	Human Testes	Testis			pCMVSport
1102.52						1
H0373	Human Heart	Human Adult Heart	Heart			pCMVSport
						1

H0383	Human Prostate BPH, re-	Human Prostate BPH				Uni-ZAP XR
H0392	H. Meningima, M1	Human Meningima	brain	+	<u> </u>	pSport1
H0399	Human Kidney Cortex, re-	Human Kidney	014			Lambda
	rescue	Cortex		1	i	ZAP II
H0411	H Female Bladder, Adult	Human Female Adult Bladder	Bladder			pSport1
H0412	Human umbilical vein	HUVE Cells	Umbilical	Cell Line		pSport1
	endothelial cells, IL-4 induced		vein			,
H0414	Ovarian Tumor I, OV5232	Ovarian Tumor, OV5232	Ovary		disease	pSport1
H0415	H. Ovarian Tumor, II, OV5232	Ovarian Tumor, OV5232	Ovary		disease	pCMVSport 2.0
H0427	Human Adipose	Human Adipose, left hiplipoma				pSport1
H0428	Human Ovary	Human Ovary Tumor	Ovary			pSport1
H0431	H. Kidney Medulla, re- excision	Kidney medulla	Kidney			pBluescript
H0435	Ovarian Turnor 10-3-95	Ovarian Tumor, OV350721	Ovary			pCMVSport
H0436	Resting T-Cell Library,Il	T-Cells	Blood	Cell Line		pSport1
H0478	Salivary Gland, Lib 2	Human Salivary Gland	Salivary gland			pSport i
H0483	Breast Cancer cell line, MDA 36	Breast Cancer Cell line, MDA 36				pSport1
H0484	Breast Cancer Cell line, angiogenic	Breast Cancer Cell line, Angiogenic, 36T3	1			pSport1
H0486	Hodgkin"s Lymphoma II	Hodgkin"s Lymphoma II			disease	pCMVSport
H0494	Keratinocyte	Keratinocyte				pCMVSport 2.0
H0520	NTERA2 + retinoic acid, 14 days	NTERA2, Teratocarcinoma cell line				pSport1
H0521	Primary Dendritic Cells, lib 1	Primary Dendritic cells	-			pCMVSport 3.0
H0533	Human Stromal endometrial fibroblasts, treated w/ estradiol	Human Stromal endometrial fibroblasts, treated wit				pSport1
H0534	Human Stromal endometrial fibroblasts, treated with progesterone	Human Stromal endometrial ' fibroblasts, treated w/				pSport1
H0539	Pancreas Islet Cell Tumor	Pancreas Islet Cell Tumour	Pancreas		disease	pSport1
H0543	T cell helper II	Helper T cell				pCMVSport 3.0
H0544	Human endometrial stromal cells	Human endometrial stromal cells				pCMVSport 3.0
H0545	Human endometrial stromal cells-treated with progesterone	Human endometrial stromal cells-treated with proge				pCMVSport 3.0
H0546	Human endometrial stromal cells-treated with estradiol	Human endometrial stromal cells-treated with estra				pCMVSport 3.0
H0547	NTERA2 teratocarcinoma cell line+retinoic acid (14 days)	NTERA2, Teratocarcinoma cell line				pSport1

H0549	H. Epididiymus, caput &	Human	T		'	Uni-ZAP XR
	corpus	Epididiymus, caput				Olli-ZAI XI
		and corpus			<u> </u>	
H0550	H. Epididiymus, cauda	Human Epididiymus, cauda				Uni-ZAP XI
H0551	Human Thymus Stromal Cells	Human Thymus Stromal Cells				pCMVSport 3.0
H0553	Human Placenta	Human Placenta				pCMVSport
H0555	Rejected Kidney, lib 4	Human Rejected Kidney	Kidney		disease	pCMVSport
H0560	КМН2	KMH2				pCMVSport
H0587	Healing groin wound; 7.5 hours post incision	Groin-2/19/97	groin	,	disease	pCMVSport
H0592	Healing groin wound - zero hr post-incision (control)	HGS wound healing , project; abdomen			disease	pCMVSport 3.0
H0593	Olfactory epithelium;nasalcavity	Olfactory epithelium from roof of left nasal cacit				pCMVSport 3.0
H0596	Human Colon Cancer;re- excision	Human Colon Cancer	Colon			Lambda ZAP II
H0606	Human Primary Breast Cancer;re-excision	Human Primary Breast Cancer	Breast		disease	Uni-ZAP XR
H0615	Human Ovarian Cancer Reexcision	Ovarian Cancer	Ovary		disease	Uni-ZAP XR
H0616	Human Testes, Reexcision	Human Testes	Testis			Uni-ZAP XR
H0617	Human Primary Breast Cancer Reexcision	Human Primary Breast Cancer	Breast '		disease	Uni-ZAP XR
H0618	Human Adult Testes, Large Inserts, Reexcision	Human Adult Testis	Testis	-		Uni-ZAP XR
H0623	Human Umbilical Vein; Reexcision	Human Umbilical Vein Endothelial Cells	Umbilical vein			Uni-ZAP XR
H0632	Hepatocellular Tumor;re- excision	Hepatocellular Turnor	Liver			Lambda ZAP II
H0634	Human Testes Tumor, re- excision	Human Testes Tumor	Testis		disease	Uni-ZAP XR
H0641	LPS activated derived dendritic cells	LPS activated monocyte derived dendritic cells	,			pSport1
H0643	Hep G2 Cells, PCR library	Hep G2 Cells				Other
Н0644	Human Placenta (re- excision)	Human Placenta	Placenta			Uni-ZAP XR
H0646	Lung, Cancer (4005313 A3): Invasive Poorly Differentiated Lung Adenocarcinoma,	Metastatic squamous cell lung carcinoma, poorly di				pSport1
H0647	Lung, Cancer (4005163 B7): Invasive, Poorly Diff. Adenocarcinoma, Metastatic	Invasive poorly differentiated lung adenocarcinoma			disease	pSportI
H0648	Ovary, Cancer: (4004562 B6) Papillary Serous Cystic Neoplasm, Low Malignant Pot	Papillary Cstic neoplasm of low malignant potentia			disease	pSport1
H0650	B-Cells	B-Cells				pCMVSport 3.0
H0651	Ovary, Normal: (9805C040R)	Normal Ovary				pSport1
H0657	B-cells (stimulated)	B-cells (stimulated)				pSport1

WO 01/55320

differentiated adenocarcinoma Grade II Papillary Covary Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Carcinoma Grade II Papillary Grade II Papillary G	H0658	Ovary, Cancer	9809C332- Poorly	Ovary &	 	disease	pSport1-
H0669			differentiate	Fallopian Tubes			
Hoffer Papillary Carcinoma Carcinoma, Ovary Papillary Carcinoma Carcinoma, Ovary Papillary Carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated carcinoma Poorly differentiated	110650		<u> </u>	<u> </u>	_		
15799A IF) Poorly differentiated carcinoma 1600522 160052282) 1600528282) 1600528282) 1600528282) 1600528282) 1600528282) 1600528282) 1600528282) 1600528282) 1600528282) 160052828282) 160052828282) 160052828282] 16005282828282] 160052828282828282828282828282828282828282	H0639	(15395A1F): Grade II		Ovary		disease	pSport1
H0662 Breast, Cancer: (4004943 Breast cancer A5 Breast Breast Breast Breast PSport1	H0660	(15799A1F) Poorly				disease	pSport1
H0663 Breast, Cancer: (4005522B2) #4005522(B2) Breast Cancer Breast disease pSport1	H0661	Breast, Cancer: (4004943	Breast cancer			disease	pSport1
H0664	H0662	Breast, Normal: (4005522B2)	Normal Breast - #4005522(B2)	Breast			pSport1
H0666	H0663		Breast Cancer -	Breast	,	disease	pSport1
A2 Sample #4004332A2	H0664			Breast		disease	pSport1
A33: Well-Differentiated Micropapillary Serous Carcinoma Cancer(4004576A8) PSport1		A2)	Sample			disease	pSport1 ·
A8 Cancer(4004576A8 Prostate		A3): Well-Differentiated Micropapillary Serous Carcinoma					pSport1
Human Prostate Cancer, Stage B2; re-excision	H0672	A8)		Ovary			pSport1
Human Prostate Cancer, Stage C Human Prostate Cancer, stage C Colon, Cancer: Colon Cancer 9808C064R 9808C064R 9808C064R Other	H0673		Human Prostate	Prostate			Uni-ZAP XR
H0676 Colon, Cancer: (9808C064R)-total RNA Placenta Placenta Placenta Other	H0674		Human Prostate	Prostate	,		Uni-ZAP XR
H0683 Screened clones from placental library Placenta Placenta Placenta Placenta Placenta Placenta Placenta Other	H0676		Colon Cancer				
Adenocarcinoma adenocarcinoma, stage 3C (9804G01 3.0	H0678	placental library	Placenta	Placenta			
Adenocarcinoma of Ovary, Human Cell Line, #OVCAR-3	H0683		adenocarcinoma,				
Ovary, Human Cell Line	H0685	Ovary, Human Cell Line,	Adenocarcinoma of Ovary, Human Cell				
Human normal ovary(#9610G215) Human normal ovary(#9610G215) Human normal ovary(#9610G215) Human Ovarian Cancer(#9807G017) Human Ovarian Cancer(#9807G017) Human Ovarian Cancer(#9807G017) Human Ovarian Cancer(#9807G017) Human Ovarian Cancer	H0686		Ovary, Human Cell				
Human Ovarian Cancer (#9807G017) Human Ovarian Cancer (#9807G017) MRNA from Maura Ru PCMVSport 3.0		ovary(#9610G215)	Human normal	Ovary			
#9806G019	H0688		cancer(#9807G017), mRNA from Maura	-			pCMVSport
H0690 Ovarian Cancer, # 9702G001 #9702G001 S0004 Prostate Prostate Prostate Prostate Lambda ZAP II S0013 Prostate Prostate prostate Uni-ZAP XR S0014 Kidney Cortex Kidney Cortex Kidney Uni-ZAP XR S0026 Stromal cell TF274 stromal cell Bone marrow Cell Line Uni-ZAP XR S0028 Smooth muscle, control Smooth muscle Pulmanary artery Cell Line Uni-ZAP XR	H0689						
S0004 Prostate Prostate Prostate Prostate Lambda ZAP II S0013 Prostate Prostate prostate Uni-ZAP XR S0014 Kidney Cortex Kidney cortex Kidney Uni-ZAP XR S0026 Stromal cell TF274 stromal cell Bone marrow Cell Line Uni-ZAP XR S0028 Smooth muscle, control Smooth muscle Pulmanary artery Cell Line Uni-ZAP XR		9702G001	Ovarian Cancer,			- •	pCMVSport
S0013 Prostate Prostate prostate Uni-ZAP XR S0014 Kidney Cortex Kidney cortex Kidney Uni-ZAP XR S0026 Stromal cell TF274 stromal cell Bone marrow Cell Line Uni-ZAP XR S0028 Smooth muscle, control Smooth muscle Pulmanary artery Cell Line Uni-ZAP XR	S0004	Prostate	Prostate BPH	Prostate			Lambda
S0014 Kidney Cortex Kidney Kidney Uni-ZAP XR	S0013	Prostate	Prostate	prostate			
S0026 Stromal cell TF274 stromal cell Bone marrow Cell Line Uni-ZAP XR S0028 Smooth muscle, control Smooth muscle Pulmanary artery Cell Line Uni-ZAP XR	S0014	Kidney Cortex	Kidney cortex		 		
S0028 Smooth muscle, control Smooth muscle Pulmanary artery Cell Line Uni-ZAP XR	S0026				Cell Line		Uni-ZAP XR
	S0028	Smooth muscle,control		Pulmanary			Uni-ZAP XR
	S0042	Testes	Human Testes				ZAP Express

PCT/US01/01339

S0044	Prostate BPH	prostate BPH	Prostate		1:2	Uni-ZAP XR
S0052	neutrophils control	human neutrophils	blood	Cell Line	discase	Uni-ZAP XR
S0112	Hypothalamus		Brain	Cen Ente		Uni-ZAP XR
S0134	Apoptotic T-cell	apoptotic cells	- Diani	Cell Line		
S0146	prostate-edited	prostate BPH	Prostate	Cen Line		Uni-ZAP XR
S0148	Normal Prostate	Prostate	prostate	 		Uni-ZAP XR
S0150		LNCAP Cell Line	Prostate	Cell Line		Uni-ZAP XR
S0152	PC3 Prostate cell line	PC3 prostate cell	TTOSIZIC	Cell Line		Uni-ZAP XR
'		line		1		Uni-ZAP XR
S0168	Prostate/LNCAP,	PC3 prostate cell	 	 		Di-
	subtraction I	line	İ	1		pBluescript
S0174	Prostate-BPH subtracted II	Human Prostate	 			pBluescript
1		ВРН	l	1		pBiuescript
S0176	Prostate, normal,	Prostate	prostate	 		Uni-ZAP XR
	subtraction I		prostate			UIII-ZAF AR
S0188	Prostate, BPH, Lib 2	Human Prostate		-	discase	pSport1
	<u> </u>	ВРН	ļ		discase	populti
S0190	Prostate BPH,Lib 2,	Human Prostate	†- 			pSport1
	subtracted	BPH				popoliti
S0212	Bone Marrow Stromal	Bone Marrow		1		pSport1
	Cell, untreated	Stromal]		poporti
		Cell,untreated	1	! !		
S0222	H. Frontal	H. Brain, Frontal	Brain		disease	Uni-ZAP XR
	cortex,epileptic;re-	Cortex, Epileptic	1			
00010	excision					
S0242	Synovial Fibroblasts	Synovial Fibroblasts				pSport1
22251	(III/TNF), subt					1 ' '
S0274	PCMIX	PCMIX (Human	Brain			PCRII
00000		Cerebellum)				
S0282	Brain Frontal Cortex, re-	Brain frontal cortex	Brain			Lambda
C0204	excision					ZAP II
S0284	7TMCTT (Testis)	7TMCTP (Placenta)	Testis			PCRII
S0286	7TMCTP (Placenta)	Н7МСТР	Placenta			PCRII
S0294	1	(PLACENTA)				
30294	Larynx tumor	Larynx tumor	Larynx,vocal		disease	pSport1
S0326	Mammary Gland	N- Cl 1	cord		·	
30320	Manufary Giand	Mammary Gland	Whole			pSport1
ĺ	1		mammary	1		'
S0328	Palate carcinoma	Palate carcinoma	gland			
S0352	Larynx Carcinoma		Uvula		disease	pSport1
S0354	Colon Normal II	Larynx carcinoma Colon Normal	0.1.		disease	pSport1
S0358	Colon Normal III	Colon Normal	Colon			pSport1
S0360	Colon Tumor II	Colon Tumor	Colon			pSport1
S0374	Normal colon	Normal colon	Colon		disease	pSport1
S0374	Pancreas Turnor PCA4 Tu	Pancreas Tumor			1.	pSport1
00300	Tancicas Tulliol FCA4 Tu	PCA4 Tu			disease	pSport1 .
S0396	Uterus; normal	Uterus; normal				
S0398	Testis; normal	Testis; normal				pSport1
S0412	Temporal cortex-	Temporal cortex,				pSport1
-0	Alzheizmer; subtracted	alzheimer	j	ļ	disease	Other
S0422	Mo7e Cell Line GM-CSF	Mo7e Cell Line				-6141/5
00.00	treated (lng/ml)	GM-CSF treated	[pCMVSport
	((lng/ml)	İ			3.0
S0424	TF-1 Cell Line GM-CSF	TF-1 Cell Line				nSnort!
	Treated	GM-CSF Treated	į			pSport1
S0442	Colon Normal	Colon Normal				nSport1
S0444	Colon Tumor	Colon Tumour			disease	pSport1
S0454	Placenta	Placenta	Placenta	 +	discase	pSport1
S0456	Tongue Normal	Tongue Normal	1 faccina			pSport1
S0460	Thyroid Tumour	Thyroid Tumour				pSport1
	,	zarya zamou				pSport1

T0010	Human Infant Brain	Human Infant Brain	1	1		Other
T0041	Jurkat T-cell G1 phase	Jurkat T-cell		-		pBluescript
)	Juntal 1 cell				,
T0068	Normal Ovary,	Normal Ovary,	 			SK-
10000	Premenopausal	Premenopausal			1	pBluescript
T0069	Human Uterus, normal			<u> </u>		SK-
10009	Human Oterus, normai	Human Uterus,			1	pBluescript
10005	 	normal				SK-
L0005	Clontech human aorta	j		ł		
	polyA+ mRNA (#6572)			1		.1
_L0021	Human adult (K.Okubo)					,
L0023	human adult testis				 	
L0040	Human colon mucosa		<u> </u>	-	 	
L0041	Human epidermal		 		-}	
	keratinocyte					
L0060	Human thymus NSTH II	 	 			
L0070	Selected chromosome 21			<u> </u>		
LOUTO			1	ŀ		
10100	cDNA library					
L0109	Human brain cDNA	brain				
L0142	Human placenta cDNA	placenta				
	(TFujiwara)			,		
L0143	Human placenta polyA+	placenta				
<u> </u>	(TFujiwara)					1
L0151	Human testis (C. De	testis				
[Smet)	1		1	1	1
L0157	Human fetal brain		brain	 	 -	
	(TFujiwara)		Ulatti			i i
L0163	Human heart cDNA		 			
L0103	(YNakamura)		heart		İ	
L0351	Infant brain, Bento Soares		<u> </u>		<u> </u>	
L L0331	Iniani orain, Bento Soares	'		1		BA, M13-
1.0250	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		1			derived
L0352	Normalized infant brain,			1		BA, M13-
<u> </u>	Bento Soares			1	Ì	derived
L0361	Stratagene ovary	,	ovary			Bluescript
	(#937217)		•			SK
L0362	Stratagene ovarian cancer					Bluescript
	(#937219)		1	i	1	SK-
L0363	NCI_CGAP_GC2	germ cell tumor				Bluescript
1		8		1		SK-
L0365	NCI_CGAP_Phe1	pheochromocytoma			 -	
		phedemoniocytoma				Bluescript
L0366	Stratagene schizo brain	schizophrenic brain	 		<u> </u>	SK-
20300	S11				İ	Bluescript
L0367	NCI_CGAP_Sch1	S-11 frontal lobe	ļ		 	SK-
L0307	NCI_CGAP_Scn1	Schwannoma tumor	ĺ	ľ		Bluescript
1 22 22	NG SC S S S S S S S S S S S S S S S S S S					SK-
L0369	NCI_CGAP_AA1	adrenal adenoma	adrenal gland			Bluescript
		•				SK-
L0371	NCI_CGAP_Br3	breast tumor	breast			Bluescript
L						SK-
L0372	NCI_CGAP_Co12	colon tumor	colon		 	Bluescript
					1	SK-
L0373	NCI_CGAP_Col1	· tumor	colon			Bluescript
	-		201011		1 .	SK-
L0374	NCI_CGAP_Co2	tumor	colon		 	
		tantoi	COION			Bluescript
L0375	NCI CGAP Kid6	In the same of the				SK-
[[[]	NCI_CGAP_KIGO	kidney tumor	kidney			Bluescript
10055	NGLOGIA					SK-
L0376	NCI_CGAP_Larl	larynx	larynx		}	Bluescript
						SK-
L0378	NCI_CGAP_Lu1	lung tumor	lung			Bluescript
					-	SK-
L0380	NCI_CGAP_HN1	squamous cell	lymph node			Bluescript
						

	1				
L0381	NCI_CGAP_HN4	, carcinoma	ļ		SK-
1.0361	NCI_COAP_HN4	squamous cell	pharynx		Bluescript
L0383	NCI_CGAP_Pr24	carcinoma	 	 	SK-
L0303	NCI_COAF_F124	invasive tumor (cell	prostate		Bluescript
L0411	I-NIB	line)	 	 	SK-
L0435	Infant brain, LLNL array		<u> </u>		Lafmid BA
1 20133	of Dr. M. Soares INIB				lafmid BA
L0438	normalized infant brain	total brain	brain	 	
	cDNA	total olalli	оган	1	lafmid BA
L0439	Soares infant brain 1NIB		whole brain	 	1.6:154
L0441	2HB3MK	· · · · · · · · · · · · · · · · · · ·	whole brain	 	Lafmid BA
L0455	Human retina cDNA	retina	eye	 -	Lafmid BK
	randomly primed	1	Cyc		lambda gt10
	sublibrary	1			
L0462	WATM1				lambda gt l l
L0465	TEST1, Human adult				lambda
	Testis tissue			1	nml 149
L0470	BL29 Burkitt"s	-			lambda ZAP
	lymphoma, Pascalis		'		2
10471	Sideras				
L0471	Human fetal heart,				Lambda
L0475	Lambda ZAP Express				ZAP Express
1.0473	KG1-a Lambda Zap Express cDNA library			KG1-a	Lambda Zap
	Express CDIVA Horary			·]	Express
L0480	Stratagene cat#937212				(Stratagene)
20.00	(1992)				Lambda
	(1222)			ļ	ZAP,
					pBluescript SK(-)
L0483	Human pancreatic islet		-	. • • • • • • • • • • • • • • • • • • •	Lambda
		1		.	ZAPII
L0485	STRATAGENE Human	skeletal muscle	leg muscle		Lambda
	skeletal muscle cDNA	,			ZAPII
10.00	library, cat. #936215.				
L0492	Human Genomic				pAMP
L0509	NCI_CGAP_Lu26	invasive	lung		pAMPI
L0512	NGL CCAR O 26	adenocarcinoma			
L0312	NCI_CGAP_Ov36	borderline ovarian	ovary	1	pAMP1
L0513	NCI_CGAP_Ov37	carcinoma early stage papillary			
20313	NCI_COAI_OV37	serous carcinoma	ovary		pAMP1
L0515	NCI_CGAP_Ov32	papillary serous			
	7.01_007.11_07.52	carcinoma	ovary	1	pAMPI
L0516	Chromosome 19p12-p13.1	Garcinoma			-AMD10
	exon				pAMP10
L0517	NCI_CGAP_Pr1				pAMP10
L0518	NCI_CGAP_Pr2				pAMP10
L0519	NCI_CGAP_Pr3				pAMP10
L0520	NCI_CGAP_Alv1	alveolar		. 	pAMP10
		rhabdomyosarcoma	i	1	PARIATI IO
L0521	NCI_CGAP_Ew1	Ewing"s sarcoma			pAMP10
L0522	NCI_CGAP_Kid1	kidney			pAMP10
L0523	NCI_CGAP_Lip2	liposarcoma			pAMP10
L0524	NCI_CGAP_Li1	liver			pAMP10
L0526	NCI_CGAP_Pri2	metastatic prostate	•		pAMP10
		bone lesion			F
L0527	NCI_CGAP_Ov2	ovary			pAMP10
L0528	NCI_CGAP_Pr5	prostate			pAMP10 ·
L0529	NCI_CGAP_Pr6	prostate			pAMP10
L0532	NCI_CGAP_Thy1	thyroid			pAMP10

WO 01/55320

L0533	NCI CGAP HSCI	stem cells	bone marrow	, , '		1 11/010
L0534	Chromosome 7 Fetal	brain	brain	' 		pAMP10
	Brain cDNA Library	1	- Cram	· '	ĺ	PAMIFIU
L0536	NCI_CGAP_Br4	normal ductal tissue	breast .			pAMP10
L0539		1	placenta			pAMP10
L0542	CDNA Library NCI_CGAP_Prl1					
10342	NCI_COMF_PIII	normal prostatic epithelial cells	prostate			pAMP10
L0547	NCI CGAP Pr16	tumor	prostate	 	+	pAMP10
L0558	NCI_CGAP_Ov40	endometrioid	ovary		-	pAMP10
L0565	111	ovarian metastasis	<u></u>		,	
10303	Normal Human Trabecular Bone Cells	Bone	Hip			pBluescript
L0581	Stratagene liver (#937224)		liver	+		pBluescript
				'		SK
L0586	HTCDL1					pBluescript
L0587	Stratagene colon HT29		 	 		SK(-)
20507	(#937221)		j			pBluescript SK-
L0588	Stratagene endothelial cell			+		pBluescript
1050	937223					SK-
L0589	Stratagene fetal retina 937202	•				pBluescript
L0590	Stratagene fibroblast		 	 		SK-
	(#937212)					pBluescript SK-
L0591	Stratagene HeLa cell s3			 	 	pBluescript
L0592	937216					SK-
L0392	Stratagene hNT neuron (#937233)			1	İ	pBluescript
L0593	Stratagene		<u> </u>	 		SK- pBluescript
	neuroepithelium	,	1	1,'		SK-
1.0504	(#937231)					
L0594	Stratagene neuroepithelium					pBluescript
	NT2RAMI 937234					SK-
L0595	Stratagene NT2 neuronal	neuroepithelial cells	brain		<u> </u>	pBluescript
1.0506	precursor 937230					SK-
L0596	Stratagene colon (#937204)		colon			pBluescript
L0597	Stratagene corneal stroma		cornea		ļ	SK-
	(#937222)		comea	1		pBluescript SK-
L0598	Morton Fetal Cochlea	cochlea	ear			pBluescript
L0599	Stratagene lung (#937210)			ļ <u>.</u>		SK-
L0399	Stratagene lung (#93/210)	•	lung			pBluescript
L0600	Weizmann Olfactory	olfactory epithelium	nose	<u> </u>		SK pBluescript
	Epithelium					SK-
L0601	Stratagene pancreas (#937208)		pancreas			pBluescript
L0602	Pancreatic Islet	pancreatic islet	nonce	 		SK-
		panereatic isiet	pancreas			pBluescript SK-
L0603	Stratagene placenta		placenta	 	 	pBluescript
10001	(#937225)		· · · · · · · · · · · · · · · · · · ·			SK-
L0604	Stratagene muscle 937209	muscle	skeletal			pBluescript
L0605	Stratagene fetal spleen	fetal spleen	muscle		 	SK-
	(#937205)	iciai spiccii	spłeeп			pBluescript SK-
L0606	NCI_CGAP_Lym5	follicular lymphoma	lymph node		<u> </u>	pBluescript
10600	Carte					SK-
L0608	Stratagene lung carcinoma 937218	lung carcinoma	lung	NCI-H69		pBluescript
				<u> </u>	L	SK-

L0612	Schiller	oligodendroglioma	brain	 		pBluescript
	oligodendroglioma	- Ingodenaregnenia	J. J. J. J. J. J. J. J. J. J. J. J. J. J			SK-
				1	Į.	(Stratagene)
L0617	Chromosome 22 exon					pBluescriptII
						KS+
L0618	Chromosome 9 exon					pBluescriptll
						KS÷
L0623	HM3	pectoral muscle				pcDNAII
		(after mastectomy)				(Invitrogen)
L0626	NCI_CGAP_GC1	bulk germ cell				pCMV-
		seminoma				SPORT2
L0635	NCI_CGAP_PNS1	dorsal root ganglion	peripheral			pCMV-
			nervous		1	SPORT4
10027	NCL COLD D. CO.		system			
L0637	NCI_CGAP_Bm53	three pooled	brain			pCMV-
1.0629	NGL CCAP P. 26	meningiomas				SPORT6
L0638	NCI_CGAP_Bm35	tumor, 5 pooled (see	brain		1	pCMV-
L0639	NCI_CGAP_Brn52	description)			<u> </u>	SPORT6
L0039	NCI_CGAP_Bm32	tumor, 5 pooled (see	brain		Ì	pCMV-
L0642	NCI_CGAP_Co18	description)	 	(- 	SPORT6
L0042	NCI_COAF_COIS	moderately differentiated	colon			pCMV-
	·	adenocarcinoma				SPORT6
L0646	NCI_CGAP_Co14	moderately-	colon		-	
20040	NCI_COAI_COI4	differentiated	colon			pCMV-
		adenocarcinoma	ļ			SPORT6
L0647	NCI_CGAP_Sar4	five pooled	connective		 	-0107
		sarcomas, including	tissue		:	pCMV- SPORT6
		myxoid liposarcoma	l 135uc			SPORTO
L0649	NCI_CGAP_GUI	2 pooled high-grade	genitourinary		+	pCMV-
	· - -	transitional cell	tract		Ì	SPORT6
	'	tumors			J	Si Okto
L0655	NCJ_CGAP_Lym12	lymphoma,	lymph node			pCMV-
		follicular mixed	, ,		1	SPORT6
		small and large cell			1	
L0656	NCI_CGAP_Ov38	normal epithelium	ovary			pCMV-
					J	SPORT6
L0657	NCI_CGAP_Ov23	tumor, 5 pooled (see	ovary			pCMV-
10650	2101 0010 0	description)			ļ	SPORT6
L0658	NCI_CGAP_Ov35	tumor, 5 pooled (see	ovary			pCMV-
L0659	NCL CCAR P.D	description)			ļ	SPORT6
L0039	NCI_CGAP_Pan1	adenocarcinoma	pancreas			pCMV-
L0661	NCI_CGAP_Mel15			 .		SPORT6
LUGGI	NCI_COAF_Mell3	malignant melanoma,	skin			pCMV-
-		metastatic to lymph				SPORT6
1		node			İ	
L0662	NCI_CGAP_Gas4	poorly differentiated	stomach		 	pCMV-
	0000	adenocarcinoma	Stomach			SPORT6
1		with signet r	[1	SICKIO
L0663	NCI_CGAP_Ut2	moderately-	uterus			pCMV-
ļ	- -	differentiated				SPORT6
İ		endometrial	1			
		adenocarcino	1		1	
L0664	NCI_CGAP_Ut3	poorly-differentiated	uterus			pCMV-
1	•	endometrial			1	SPORT6
		adenocarcinoma,				
L0665	NCI_CGAP_Ut4	serous papillary	uterus			pCMV-
İ		carcinoma, high	į			SPORT6
		grade, 2 pooled t				<u> </u>
L0666	NCI_CGAP_Ut1 .	well-differentiated	uterus			pCMV-
- 1		endometrial	Į.		1	SPORT6

1000	NCL CCAR COM	adenocarcinoma, 7				<u> </u>
L0667	NCI_CGAP_CML1	myeloid cells, 18	whole blood	1		pCMV-
	1	pooled CML cases,	1.		1	SPORT6
<u> </u>		BCR/ABL rearra	<u> </u>		i	İ
L0686	Stanley Frontal SN pool 2	frontal lobe (see	brain			pCR2.1-
		description)		i		TOPO
			f	Í	1	(Invitrogen)
L0697	Testis I			 	 	PGEM
L'				l		5zf(+)
L0698	Testis 2			 		PGEM
L0700	Outward Alu-primed	·				5zf(+)
	hncDNA library			1		pGEM-3Z
L0717	Gessler Wilms tumor					
L0731	Soares_pregnant_uterus_	· · · · · · · · · · · · · · · · · · ·		ļ	 	pSPORT1
L0/31	NbHPU		uterus			pT7T3-Pac
L0738						
	Human colorectal cancer					pT7T3D
L0740	Soares melanocyte	melanocyte		i		pT7T3D
ŀ	2NbHM	, ,				(Pharmacia)
		· I		1 ,	1	with a
						modified
ļ.,						polylinker
L0741	Soares adult brain		brain			pT7T3D
	N2b4HB55Y					(Pharmacia)
1	1		•		1	with a
					f	modified
]			1	ł	polylinker
L0742	Soares adult brain		brain		†	pT7T3D
i	N2b5HB55Y		0.4			(Pharmacia)
i		•				with a
			ı	-		
1	'				1	modified
L0743	Soares breast 2NbHBst		breast		 	polylinker
	Source Breast 2, 10,113,		breast		1	pT7T3D
					1	(Pharmacia)
					1	with a
					i	modified
L0744	Soares breast 3NbHBst					polylinker
LUTT	Soales Oleast SNOTIDSt		breast]	pT7T3D
		•			1	(Pharmacia)
						with a
			i		1	modified
1.0745	C					polylinker
L0745	Soares retina N2b4HR	retina	eye		1	pT7T3D
		ľ				(Pharmacia)
	- 1	1				with a
			•			modified
105:1						polylinker
L0746	Soares retina N2b5HR	retina	eye			pT7T3D
	!				1	(Pharmacia)
	1		ļ			with a
	1		i		1	modified
			i			polylinker
L0747	Soares_fetal_heart_NbHH		heart			pT7T3D
	19W		l			(Pharmacia)
	i		l			with a
	ŀ		ĺ			modified
		1	ļ			
L0748	Soares fetal liver spleen		Liver and			polylinker
	INFLS	İ				pT7T3D
		l l	Spleen			(Pharmacia)
ŀ		İ			-	with a
İ	J		j			modified
			L		L	polylinker

PCT/US01/01339

L0749		1. 1	Liver and			pT7T3D
	_INFLS_SI		Spleen	1 ,	1	(Pharmacia)
			,	1.	1	with a
	•			·		modified
		1 ' .	·	ŀ		polylinker
L0750	Soares_fetal_lung NbHL1		lung	1	 	pT7T3D
1	19W		15	1	ł	(Pharmacia)
						with a
		İ	1	1]	modified
	· ·			1	İ	
L0751	Soares ovary tumor	ovarian tumor	0110711	 	 	polylinker
	NbHOT	Ovarian tunior	ovary	1		pT7T3D
	1.00.00			1	1	(Pharmacia)
			ļ	1	[with a
	1					modified
L0752	Soares_parathyroid_tumor		 	'	 	polylinker
L0/32	NbHPA	parathyroid tumor	parathyroid		ł	pT7T3D
	_NORFA		gland			(Pharmacia)
1					Į.	with a
						modified ,
1.0753		<u> </u>		1		połylinker
L0753	Soares_pineal_gland_N3H		pineal gland			pT7T3D
	PG			1		(Pharmacia)
İ				İ		with a
			ļ		1	modified
						polylinker
L0754	Soares placenta Nb2HP		placenta			pT7T3D
			1 .			(Pharmacia)
-	•				1	with a
				ŀ		modified
L			1 '	l		polylinker
L0755	Soares_placenta_8to9wee		placenta '	i i		pT7T3D
	ks_2NbHP8to9W		Piaseilla	,		(Pharmacia)
İ	-	•	ĺ	1 .	1	with a
	·					modified
1				Ì	ļ	polylinker.
L0756	Soares_multiple_sclerosis	multiple sclerosis				pT7T3D
	_2NbHMSP	lesions				
1	_	100.0115	ļ	1		(Pharmacia) with a
ł	İ			1	[1
	1		1			modified
1			1	1		polylinker
L0757	Soares_senescent_fibrobla	senescent fibroblast	ļ			V TYPE
20737	sts_NbHSF	sellescelli libioblasi			1	pT7T3D
	313_1101131	*	ł			(Pharmacia)
	i i				l	with a
						modified
					Į.	polylinker
L0758	Soares_testis_NHT				 	V_TYPE
1 20/38	Somes Testis INLI				1	pT7T3D-Pac
1			1		1	(Pharmacia)
1					1	with a
	'					modified
10770	Canada and Canada					polylinker
L0759	Soares_total_fetus_Nb2H		l i			pT7T3D-Pac
1	F8_9w					(Pharmacia)
1	· •					with a
	ļ					modified
ļ						polylinker
L0761	NCI_CGAP_CLL1	B-cell, chronic				pT7T3D-Pac
	ļ	lymphotic leukemia				(Pharmacia)
			ļ			with a
						modified
						polylinker
L0762	NCI_CGAP_Br1.1	breast	_			pT7T3D-Pac
			<u> </u>			P171313-1 40

						
İ						(Pharmacia)
			İ			with a
1				į	1	modified
L0763	NCI_CGAP_Br2		.			polylinker
1 20.03	Nei_eGAI_BI2	breast		Ì		pT7T3D-Pac
						(Pharmacia)
				l l		with a
1	1	j		ľ		modified
L0764	NCI_CGAP_Co3	colon				polylinker
	1	Colon				pT7T3D-Pac
		•				(Pharmacia) with a
1	ŀ	1			j	modified
						polylinker
L0765	NCI_CGAP_Co4	colon			 	pT7T3D-Pac
		1	1			(Pharmacia)
				i		with a
				İ		modified
			İ		1	polylinker
L0766	NCI_CGAP_GCB1	germinal center B				pT7T3D-Pac
		cell			•	(Pharmacia)
İ				ŀ	1	with a
						modified
1.05.5					<u>L</u> _	polylinker
L0767	NCI_CGAP_GC3	pooled germ cell				pT7T3D-Pac
İ	į.	tumors		1	i	(Pharmacia)
		1				with a
						modified
L0768	NCI_CGAP_GC4			<u> </u>		polylinker
L0708	NCI_COAP_GC4	pooled germ cell	t			pT7T3D-Pac
		tumors				(Pharmacia)
	ł			1	l	with a
ļ						modified
L0769	NCI_CGAP_Brn25	anaplastic	brain		ļ	polylinker
		oligodendroglioma	Diani	1		pT7T3D-Pac (Pharmacia)
	i	ongouding inchia			•	with a
ļ						modified
]		1	1	polylinker
L0770	NCl_CGAP_Brn23	glioblastoma	brain			pT7T3D-Pac
		(pooled)		İ		(Pharmacia)
					ļ	with a
						modified
						polylinker
L0771	NCI_CGAP_Co8	adenocarcinoma	colon			pT7T3D-Pac
		,			l	(Pharmacia)
				1	ĺ	with a
						modified
L0772	NCI_CGAP_Co10			<u> </u>		polylinker
L0772	NCI_CGAP_C010	colon tumor RER+	colon			pT7T3D-Pac
				ŀ		(Pharmacia)
				1		with a
				1		modified
L0773	NCI_CGAP_Co9	colon tumor RER+	colon	 		polylinker
23,73		COION IUNION RERT	colon	1		pT7T3D-Pac
						(Pharmacia) with a
						modified
	•					polylinker
L0774	NCI_CGAP_Kid3		kidney	 		pT7T3D-Pac
				1		(Pharmacia)
						with a
						modified
						

	<u> </u>	 	T	Τ		polylinker
L0775	NCI_CGAP_Kid5	2 pooled tumors	kidney	 -		pT7T3D-Pac
LUTTS	NCI_CGAP_KIGS		Kidney		1	
]	(clear cell type)			+	(Pharmacia)
-						with a
						modified
	<u> </u>				L	polylinker
L0776	NCI_CGAP_Lu5	carcinoid	lung	1		pT7T3D-Pac
1			}	1		(Pharmacia)
	1			1	1	with a
		ĺ	l '	1		modified
				!	1.	polylinker
L0777	Soares_NhHMPu_S1	Pooled human	mixed (see			pT7T3D-Pac
20	000.05_/ 11 0_0/	melanocyte, fetal	below)			(Pharmacia)
		heart, and pregnant	(CCIOW)		1	with a
		licart, and pregnant	1		l l	modified
					1	
1.0770	C. NEL T ODG 61		 	 	 	polylinker
L0779	Soares_NFL_T_GBC_S1		pooled			pT7T3D-Pac
		+				(Pharmacia)
		<u>,</u>	1	Ì	1	with a
		· ` `			1 1	modified
		· · · · · · · · · · · · · · · · · · ·	1			polylinker
L0780	Soares_NSF_F8_9W_OT		pooled			pT7T3D-Pac
	_PA_P_S1	ļ		1	ļ ļ	(Pharmacia)
•		İ				with a
				1	1 1	modified
			l	ŀ	!	polylinker
L0782	NCI_CGAP_Pr21	normal prostate	prostate	 	 	pT7T3D-Pac
		l	p.ostato		1	(Pharmacia)
	}			1	1	with a
	1		•	l	1	modified
	l .		F			
L0783	NGI CCAR P-22				 	polylinker_
L0/83	NCI_CGAP_Pr22	normal prostate	prostate	i	1 1	pT7T3D-Pac
	\	ļ .	ļ	Į	}	(Pharmacia)
]]	with a
						modified
						polylinker
L0786	Soares_NbHFB		whole brain	ŀ	1	pT7T3D-Pac
	l		ļ]		(Pharmacia)
				1		with a
	ł					modified
	<u> </u>]	Ì	l	1	polylinker
L0787	NCI_CGAP_Sub1		,			pT7T3D-Pac
				ļ		(Pharmacia)
•	1			ł.	1	with a
		İ		1	1	modified
	ł	'		1	, ,	polylinker
L0788	NCI_CGAP_Sub2	······		 	 	pT7T3D-Pac
D0700	1.C1_CCA1_SU02		[1	1 1	(Pharmacia)
	İ			1	į l	with a
	1	1)	1]	modified
			1	1	; l	
					 	polylinker
L0789	NCI_CGAP_Sub3	İ	1	1] [pT7T3D-Pac
				1	; l	(Pharmacia)
		ļ	1	1	Į į	with a
		j			i 1	modified
	(1		l	{	polylinker
L0790	NCI_CGAP_Sub4				,	pT7T3D-Pac
20,70			l	l	[(Pharmacia)
	1			1]]	with a
			1	1	1 1	modified
]			[
1.0501	NO. COAR C. L.S.			 	 	polylinker
L0791	NCI_CGAP_Sub5			l] [pT7T3D-Pac
	<u> </u>	L	L	L	<u> </u>	(Pharmacia)

				1		
	·	↓ . ¹		1		with a
						modified
1.0700	13101 0 0 0 0 0	<u> </u>			_	polylinker
L0792	NCI_CGAP_Sub6	,		'		pT7T3D-Pac
		1				(Pharmacia)
	j	· ·		1	}	with a
1	:			[modified
				1	1	polylinker
L0794	NCI_CGAP_GC6	pooled germ cell				pT7T3D-Pac
	,	tumors				(Pharmacia)
	1	İ '	1			with a
	14			1	ı	modified
						polylinker
L0796	NCI_CGAP_Bm50	medulloblastoma	brain		<u> </u>	pT7T3D-Pac
}	,		1	1	1	(Pharmacia)
	,					with a
		'		1	1	modified
<u></u>					İ	polylinker
L0800	NCI_CGAP_Co16	colon tumor, RER+	colon			pT7T3D-Pac
	ł		1	.		(Pharmacia)
						with a
				İ	l	modified
			L			polylinker
L0803	NCI_CGAP_Kid11		kidney		<u> </u>	pT7T3D-Pac
						(Pharmacia)
ł			1	İ		with a
						modified
L			<u></u>	1	ļ	polylinker
L0804	NCI_CGAP_Kid12	2 pooled tumors	kidney .			pT7T3D-Pac
	[(clear cell type)		1		(Pharmacia)
			,	i '		with a
J]	1		1		modified
				1 .		polylinker
L0805	NCI_CGAP_Lu24	carcinoid	lung			pT7T3D-Pac
						(Pharmacia)
						with a
						modified
10006	3101 0					polylinker
L0806	NCI_CGAP_Lu19	squamous cell	lung			pT7T3D-Pac
		carcinoma, poorly				(Pharmacia)
	1	differentiated (4		1	' i	with a
						modified
1 0007	NO. 001 0 10			L		polylinker
L0807	NCI_CGAP_Ov18	fibrotheoma	ovary			pT7T3D-Pac
-	ľ				}	(Pharmacia)
İ]		with a
	}			[1	modified
L0809	NCI CCAR P 20			ł		polylinker
FOODS	NCI_CGAP_Pr28		prostate			pT7T3D-Pac
	[1			l	(Pharmacia)
İ	1					with a
ļ				[ł	modified
!				<u> </u>	j	polylinker

TABLE 5

OMIM	Description
Reference	Description
102200	Comptetrophingue
102200	Somatotrophinoma Melo infortility due to consider de Seine
102480	Male infertility due to acrosin deficiency
	Myoadenylate deaminase deficiency
103050	Autism, succinylpurinemic
103050	Adenylosuccinase deficiency
104770	Amyloidosis, secondary, susceptibility to
106100	Angioedema, hereditary
106150	Hypertension, essential, susceptibility to
106150	Preeclampsia, susceptibility to
106300	Ankylosing spondylitis
107670	Apolipoprotein A-II deficiency
107741	Hyperlipoproteinemia, type III
107910	Virilization, maternal and fetal, from placental aromatase
	deficiency
107910	Gynecomastia, familial, due to increased aromatase activity
108725	Atherosclerosis, susceptibility to
108800	Atrial septal defect, secundum type
108962	Hypertension, salt-resistant
109270	Renal tubular acidosis, distal, 179800
109270	Spherocytosis, hereditary
109270	[Acanthocytosis, one form]
109270	[Elliptocytosis, Malaysian-Melanesian type]
109270	Hemolytic anemia due to band 3 defect
109400	Basal cell nevus syndrome
109700	Hemodialysis-related amyloidosis
110700	Vivax malaria, susceptibility to
113100	Brachydactyly, type C
113705	Ovarian cancer
113705	Breast cancer-1
113721	Breast cancer
113900	Heart block, progressive familial, type I
114240	Muscular dystrophy, limb-girdle, type 2A, 253600
116806	Colorectal cancer
118485	Polycystic ovary syndrome with hyperandrogenemia
118504	Epilepsy, benign neonatal, type 1, 121200
118504	Epilepsy, nocturnal frontal lobe, 600513
118800	Choreoathetosis, familial paroxysmal
120110	Metaphyseal chondrodysplasia, Schmid type
120120	Epidermolysis bullosa dystrophica, dominant, 131750
120120	Epidermolysis bullosa dystrophica, dolimani, 131730 Epidermolysis bullosa dystrophica, recessive, 226600
120120	Epidermolysis bullosa dystrophica, recessive, 220000 Epidermolysis bullosa, pretibial, 131850
120120	
120213	Ehlers-Danlos syndrome, type I, 130000

120216	Film Darland I W 100000
120215	Ehlers-Danlos syndrome, type II, 130010
120280	Stickler syndrome, type III
120280	Marshall syndrome, 154780
120290	OSMED syndrome, 215150
120290	Stickler syndrome, type II, 184840
120435	Muir-Torre syndrome, 158320
120435	Colorectal cancer, hereditary, nonpolyposis, type 1 Ovarian cancer
120436	Muir-Torre family cancer syndrome, 158320
120436	Turcot syndrome with glioblastoma, 276300
120436	Colorectal cancer, hereditary nonpolyposis, type 2
120700	C3 deficiency
120810	C4 deficiency
120820	C4 deficiency
120940	C9 deficiency
121011	Deafness, autosomal dominant 3, 601544
121011	Deafness, autosomal recessive 1, 220290
121014	Heterotaxia, visceroatrial, autosomal recessive
122720	Nicotine addiction, protection from
122720	Coumarin resistance, 122700
123620	Cataract, cerulean, type 2, 601547
123660	Cataract, Coppock-like
124030	Parkinsonism, susceptibility to
124030	Debrisoquine sensitivity
124200	Darier disease (keratosis follicularis)
125270	Porphyria, acute hepatic
125270	Lead poisoning, susceptibility to
125660	Myopathy, desminopathic
125660	Cardiomyopathy
125852	Insulin-dependent diabetes mellitus-2
126340	Xeroderma pigmentosum, group D, 278730
126391	DNA ligase I deficiency
126452	Autonomic nervous system dysfunction
126452	[Novelty seeking personality]
126600	Drusen, radial, autosomal dominant
126650	Chloride diarrhea, congenital, Finnish type, 214700
126650	Colon cancer
128100	Dystonia-1, torsion
129500	Ectodermal dysplasia, hidrotic
130410	Glutaricaciduria, type IIB
131100	Multiple endocrine neoplasia I
131100	Prolactinoma, hyperparathyroidism, carcinoid syndrome
131100	Carcinoid tumor of lung
131242	Shah-Waardenburg syndrome, 277580
132800	Basal cell carcinoma
132800	Epithelioma, self-healing, squamous 1, Ferguson-Smith type
172000	Department of the series of th

133171	[Frathropytosis for III-II 122100
133450	[Erythrocytosis, familial], 133100
133450	Neuroepithelioma
133701	Ewing sarcoma
	Exostoses, multiple, type 2
133780	Vitreoretinopathy, exudative, familial
134580	Factor XIIIB deficiency
134790	Hyperferritinemia-cataract syndrome, 600886
134797	Shprintzen-Goldberg syndrome, 182212
134797	Ectopia lentis, isolated
134797	Marfan syndrome, 154700
135300	Fibromatosis, gingival
135940	Ichthyosis vulgaris, 146700
136350	Pfeiffer syndrome, 101600
136435	Ovarian dysgenesis, hypergonadotropic, with normal karyotype, 233300
136550	Macular dystrophy, North Carolina type
136836	Fucosyltransferase-6 deficiency
137350	Amyloidosis, Finnish type, 105120
138079	Hyperinsulinism, familial, 602485
138079	MODY, type 2, 125851
138320	Hemolytic anemia due to glutathione peroxidase deficiency
138570	Non-insulin dependent diabetes mellitus, susceptibility to
138720	Bernard-Soulier syndrome, type B
138981	Pulmonary alveolar proteinosis, 265120
139191	Growth hormone deficient dwarfism
139320	Pituitary ACTH secreting adenoma
139320	Pseudohypoparathyroidism, type Ia, 103580
139320	Somatotrophinoma
139320	McCune-Albright polyostotic fibrous dysplasia, 174800
141750	Alpha-thalassemia/mental retardation syndrome, type 1
141800	Methemoglobinemias, alpha-
141800	Thalassemias, alpha-
141800	Erythremias, alpha-
141800	Heinz body anemias, alpha-
141850	Thalassemia, alpha-
141850	Erythrocytosis
141850	Heinz body anemia
141850	Hemoglobin H disease
141850	Hypochromic microcytic anemia
141900	Methemoglobinemias, beta-
141900	Sickle cell anemia
141900	Thalassemias, beta-
141900	Erythremias, beta-
141900	HPFH, deletion type
141900	Heinz body anemias, beta-
	Alonas oody anomias, ucid-

WO 01/55320

142000	Thalassemia due to Hb Lepore
142000	Thalassemia, delta-
142200	**************************************
142250	HPFH, nondeletion type A
	HPFH, nondeletion type G
142270	Hereditary persistence of fetal hemoglobin
142470	[Hereditary persistence of fetal hemoglobin, heterocellular]
142857	Pemphigoid, susceptibility to
142858	Beryllium disease, chronic, susceptibility to
142959	Hand-foot-uterus syndrome, 140000
143890	Hypercholesterolemia, familial
144200	Epidermolytic palmoplantar keratoderma
145001	Hyperparathyroidism-jaw tumor syndrome
145260	Pseudohypoaldosteronism, type II
145410	Opitz G syndrome, type II
145981	Hypocalciuric hypercalcemia, type II
146760	[IgG receptor I, phagocytic, familial deficiency of]
146790	Lupus nephritis, susceptibility to
147050	Atopy
147141	Leukemia, acute lymphoblastic
147200	[Kappa light chain deficiency]
147440	Growth retardation with deafness and mental retardation
147670	Rabson-Mendenhall syndrome
147670	Diabetes mellitus, insulin-resistant, with acanthosis nigricans
147670	Leprechaunism
148065	White sponge nevus, 193900
148066	Epidermolysis bullosa simplex, Koebner, Dowling-Meara, and
	Weber-Cockayne types, 131900, 131760, 131800
148066	Epidermolysis bullosa simplex, recessive, 601001
148067	Nonepidermolytic palmoplantar keratoderma, 600962
148067	Pachyonychia congenita, Jadassohn-Lewandowsky type, 167200
148069	Pachyonychia congenita, Jackson-Lawler type, 167210
148080	Epidermolytic hyperkeratosis, 113800
150270	Laryngeal adductor paralysis
151400	Leukemia/lymphoma, B-cell, 1
151440	Leukemia, T-cell acute lymphoblastoid
151670	Hepatic lipase deficiency
152200	Coronary artery disease, susceptibility to
152445	Vohwinkel syndrome, 124500
152445	Erythrokeratoderma, progressive symmetric, 602036
152760	Hypogonadotropic hypogonadism due to GNRH deficiency,
	227200
152790	Precocious puberty, male, 176410
152790	Leydig cell hypoplasia
1.50.500	36 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
153700 153880	Macular dystrophy, vitelliform type Macular dystrophy, dominant cystoid

154275	
154275	Malignant hyperthermia susceptibility 2
154276	Malignant hyperthermia susceptibility 3
155555	[Red hair/fair skin]
155555	UV-induced skin damage, vulnerability to
156225	Muscular dystrophy, congenital merosin-deficient
156850	Cataract, congenital, with microphthalmia
157170	Holoprosencephaly-2
158590	Spinal muscular atrophy-4
159001	Muscular dystrophy, limb-girdle, type 1B
160781	Cardiomyopathy, hypertrophic, mid-left ventricular chamber type
160900	Myotonic dystrophy
161015	Mitochondrial complex I deficiency, 252010
163950	Noonan syndrome-1
163950	Cardiofaciocutaneous syndrome, 115150
164009	Leukemia, acute promyelocytic, NUMA/RARA type
164200	Oculodentodigital dysplasia
164200	Syndactyly, type III, 186100
164731	Ovarian carcinoma, 167000
164953	Liposarcoma
166600	Osteopetrosis, AD, type II
167000	Ovarian cancer, serous
167250	Paget disease of bone
168461	Multiple myeloma, 254250
168461	Parathyroid adenomatosis 1
168461	Centrocytic lymphoma
168468	Metaphyseal chondrodysplasia, Murk Jansen type, 156400
168500	Parietal foramina
168610	Parkinsonism-dementia with pallidopontonigral degeneration
170261	Bare lymphocyte syndrome, type I, due to TAP2 deficiency
170995	Zellweger syndrome-2
171190	Hypertension, essential, 145500
171650	Lysosomal acid phosphatase deficiency
172400	Hemolytic anemia due to glucosephosphate isomerase deficiency
172400	Hydrops fetalis, one form
173360	Thrombophilia due to excessive plasminogen activator inhibitor
173360	Hemorrhagic diathesis due to PAI1 deficiency
173370	Plasminogen activator deficiency
173850	Polio, susceptibility to
173870	Xeroderma pigmentosum
173870	Fanconi anemia
174000	Medullary cystic kidney disease, AD
176705	Breast cancer, sporadic
176730	Diabetes mellitus, rare form
176730	Hyperproinsulinemia, familial
176730	MODY, one form
	

176930	· Dunathantia
176930	Dysprothrombinemia
177900	Hypoprothrombinemia
	Psoriasis susceptibility-1
178640	Pulmonary alveolar proteinosis, congenital, 265120
179450	Ragweed sensitivity
179605	Retinitis pigmentosa, digenic
179605	Retinitis pigmentosa-7, peripherin-related
179605	Retinitis punctata albescens
179605	Butterfly dystrophy, retinal
179605	Macular dystrophy
179755	Renal cell carcinoma, papillary, 1
180020	Retinal cone dystrophy-1
180100	Retinitis pigmentosa-1
180104	Retinitis pigmentosa-9
180297	Anemia, hemolytic, Rh-null, suppressor type, 268150
180721	Retinitis pigmentosa, digenic
180840	Susceptibility to IDDM
180901	Malignant hyperthermia susceptibility 1, 145600
180901	Central core disease, 117000
181430	Scapuloperoneal syndrome, myopathic type
182280	Small-cell cancer of lung
182380	Glucose/galactose malabsorption
182601	Spastic paraplegia-4
182860	Pyropoikilocytosis
182860	Spherocytosis, recessive
182860	Elliptocytosis-2
182900	Spherocytosis-2
185430	Atherosclerosis, susceptibility to
185800	Symphalangism, proximal
186580	Arthrocutaneouveal granulomatosis
186855	Leukemia-2, T-cell acute lymphoblastic
188070	Bleeding disorder due to defective thromboxane A2 receptor
188540	Hypothyroidism, nongoitrous
188826	Sorsby fundus dystrophy, 136900
189800	Preeclampsia/eclampsia
190020	Bladder cancer, 109800
190040	Dermatofibrosarcoma protuberans
190040	Giant-cell fibroblastoma
190040	Meningioma, SIS-related
190182	Colon cancer
190182	Colorectal cancer, familial nonpolyposis, type 6
190198	Leukemia, T-cell acute lymphoblastic
191092	Tuberous sclerosis-2
191100	Tuberous sclerosis-1
191170	Colorectal cancer, 114500
	1

191170	I : Farmeria and and
1911/0	Li-Fraumeni syndrome
	Cervical carcinoma
191290	Segawa syndrome, recessive
191315	Insensitivity to pain, congenital, with anhidrosis, 256800
192500	Jervell and Lange-Nielsen syndrome, 220400
192500	Long QT syndrome-1
193235	Vitreoretinopathy, neovascular inflammatory
193500	Rhabdomyosarcoma, alveolar, 268220
193500	Waardenburg syndrome, type I
193500	Waardenburg syndrome, type III, 148820
193500	Craniofacial-deafness-hand syndrome, 122880
194071	Wilms tumor, type 2
194071	Adrenocortical carcinoma, hereditary, 202300
200350	Acetyl-CoA carboxylase deficiency
201460	Acyl-CoA dehydrogenase, long chain, deficiency of
201910	Adrenal hyperplasia, congenital, due to 21-hydroxylase deficiency
204500	Ceroid-lipofuscinosis, neuronal 2, classic late infantile
205100	Amyotrophic lateral sclerosis, juvenile
207750	Hyperlipoproteinemia, type Ib
209901	Bardet-Biedl syndrome 1
215700	Citrullinemia
216900	Achromatopsia
217000	C2 deficiency
217050	C6 deficiency
217050	Combined C6/C7 deficiency
217070	C7 deficiency
221770	Polycystic lipomembranous osteodysplasia with sclerosing
	leukencephalopathy
221820	Gliosis, familial progressive subcortical
222100	Diabetes mellitus, insulin-dependent-1
223360	Dopamine-beta-hydroxylase deficiency
223900	'Dysautonomia, familial
227646	Fanconi anemia, type D
227650	Fanconi anemia, type A
230400	Galactosemia
230450	Hemolytic anemia due to gamma-glutamylcysteine synthetase
	deficiency
230800	Gaucher disease
230800	Gaucher disease with cardiovascular calcification
231670	Glutaricaciduria, type I
231680	Glutaricaciduria, type IIA
231950	Glutathioninuria
232200	Glycogen storage disease I
232400	Glycogen storage disease IIIa
232400	Glycogen storage disease IIIb

232600	McArdle disease
232800	
	Glycogen storage disease VII
233100	[Renal glucosuria]
235200	Hemochromatosis
237300	Carbamoylphosphate synthetase I deficiency
239500	Hyperprolinemia, type I
245050	Ketoacidosis due to SCOT deficiency
247200	Miller-Dieker lissencephaly syndrome
248600	Maple syrup urine disease, type Ia
248611	Maple syrup urine disease, type Ib
249000	Meckel syndrome
250250	Cartilage-hair hypoplasia
251170	Mevalonicaciduria
252920	Sanfilippo syndrome, type B
253000	Mucopolysaccharidosis IVA
253250	Mulibrey nanism
253700	Muscular dystrophy, limb-girdle, type 2C
253800	Walker-Warburg syndrome, 236670
253800	Fukuyama type congenital muscular dystrophy
256540	Galactosialidosis
256550	Sialidosis, type I
256550	Sialidosis, type II
256850	Giant axonal neuropathy-1
258501	3-methylglutaconicaciduria, type III
259700	Osteopetrosis, recessive
259770	Osteoporosis-pseudoglioma syndrome
261510	Pseudo-Zellweger syndrome
262000	Bjornstad syndrome
263200	Polycystic kidney disease, autosomal recessive
266200	Anemia, hemolytic, due to PK deficiency
268900	[Sarcosinemia]
270800	Spastic paraplegia-5A
272800	Tay-Sachs disease
272800	[Hex A pseudodeficiency]
272800	GM2-gangliosidosis, juvenile, adult
275350	Transcobalamin II deficiency
276700	Tyrosinemia, type I
276710	Tyrosinemia, type III
277700	Werner syndrome
278300	Xanthinuria, type I
278700	Xeroderma pigmentosum, group A
300000	Opitz G syndrome, type I
300066	Deafness, X-linked 6, sensorineural
300067	Subcortical laminar heterotopia, X-linked dominant
300067	Lissencephaly, X-linked
200001	Dissencephary, A-mikeu

300077	Mental retardation, X-linked 29
300121	
300121	Subcortical laminal heteropia, X-linked, 300067
300121	Lissencephaly, X-linked, 300067
300123	Mental retardation with isolated growth hormone deficiency
	Agammaglobulinemia, type 2, X-linked
300500	Ocular albinism, Nettleship-Falls type
300650	Ocular albinism with sensorineural deafness
301200	Amelogenesis imperfecta
301201	Amelogenesis imperfecta-3, hypoplastic type
301220	Partington syndrome II
301835	Arts syndrome
301845	Bazex syndrome
301900	Borjeson-Forssman-Lehmann syndrome
302350	Nance-Horan syndrome
302950	Chondrodysplasia punctata, X-linked recessive, 302940
304050	Aicardi syndrome
304110	Craniofrontonasal dysplasia
304340	Mental retardation, X-linked, syndromic-5, with Dandy-Walker
•	malformation, basal ganglia disease, and seizures
306100	Gonadal dysgenesis, XY female type
307150	Hypertrichosis, congenital generalized
307700	Hypoparathyroidism, X-linked
308000	HPRT-related gout
308000	Lesch-Nyhan syndrome
308700	Kallmann syndrome
309000	Lowe syndrome
309530	Mental retardation, X-linked 1, non-dysmorphic
309585	Mental retardation, X-linked, syndromic-6, with gynecomastia and
	obesity
310490	Cowchock syndrome
311200	Oral-facial-digital syndrome 1
311850	Phosphoribosyl pyrophosphate synthetase-related gout
312040	N syndrome, 310465
313850	Thoracoabdominal syndrome
600040	Colorectal cancer
600045	Xeroderma pigmentosum, group E, subtype 2
600059	Retinitis pigmentosa-13
600105	Retinitis pigmentosa-12, autosomal recessive
600119	Muscular dystrophy, Duchenne-like, type 2
600119	Adhalinopathy, primary
600140	Rubenstein-Taybi syndrome, 180849
600163	Long QT syndrome-3
600175	Spinal muscular atrophy, congenital nonprogressive, of lower limbs
600202	Dyslexia, specific, 2
600234	HMG-CoA synthease-2 deficiency
000237	THATO-COPY SAUTHERSE-7 REHICIENCY

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600261	Ehlers-Danlos-like syndrome
600266	Resistance/susceptibility to TB, etc.
600273	Polycystic kidney disease, infantile severe, with tuberous sclerosis
600276	Cerebral arteriopathy with subcortical infarcts and
(0000	leukoencephalopathy, 125310
600281	Non-insulin-dependent diabetes mellitus, 125853
600281	MODY, type 1, 125850
600309	Atrioventricular canal defect-1
600319	Diabetes mellitus, insulin-dependent, 4
600320	Insulin-dependent diabetes mellitus-5
600364	Cone dystrophy-3, 602093
600374	Bardet-Biedl syndrome 4
600528	CPT deficiency, hepatic, type I, 255120
600617	Lipoid adrenal hyperplasia, 201710
600623	Prostate cancer, 176807
600759	Alzheimer disease-4
600808	Enuresis, nocturnal, 2
600811	Xeroderma pigmentosum, group E, DDB-negative subtype, 278740
600837	Hirschsprung disease, 142623
600839	Bartter syndrome, 241200
600850	Schizophrenia disorder-4
600856	Beckwith-Wiedemann syndrome, 130650
600883	Diabetes mellitus, insulin-dependent, 8
600897	Cataract, zonular pulverulent-1, 116200
600918	Cystinuria, type III
600946	Short stature, autosomal dominant, with normal serum growth
	hormone binding protein
600946	Short stature, idiopathic
600946	Laron dwarfism, 262500
600957	Persistent Mullerian duct syndrome, type I, 261550
600958	Cardiomyopathy, familial hypertrophic, 4, 115197
600964	Refsum disease, adult, with increased pipecolicacidemia
600983	Pseudohypoaldosteronism type I, autosomal dominant, 177735
600994	Deafness, autosomal dominant 5
600996	Arrhythmogenic right ventricular dysplasia-2
601071	Deafness, autosomal recessive 9
601105	Pycnodysostosis, 265800
601154	Cardiomyopathy, dilated, 1E
601238	Cerebellar ataxia, Cayman type
601277	Ichthyosis, lamellar, type 2
601284	Hereditary hemorrhagic telangiectasia-2, 600376
601313	Polycystic kidney disease, adult type I, 173900
601316	Deafness, autosomal dominant 10
601362	DiGeorge syndrome/velocardiofacial syndrome complex-2
601363	Wilms tumor, type 4

601412 Deafness, autosomal dominant 7 601414 Retinitis pigmentosa-18 601498 Peroxisomal biogenesis disorder, complementation group 4 601517 Spinocerebellar ataxia-2, 183090 601545 Lissencephaly-1 601649 Blepharophimosis, epicanthus inversus, and ptosis, type 2 601652 Glaucoma 1A, primary open angle, juvenile-onset, 137750 601666 Insulin-dependent diabetes mellitus-15 601669 Hirschsprung disease, one form 601680 Distal arthrogryposis, type 2B 601690 Platelet-activating factor acetylhydrolase deficiency 601691 Retinitis pigmentosa-19, 601718 601691 Stargardt disease-1, 248200 601691 Cone-rod dystrophy 3 601691 Fundus flavimaculatus with macular dystrophy, 248200 601718 Retinitis pigmentosa-19 601744 Systemic lupus erythematosus, susceptibility to, 1 601757 Rhizomelic chondrodysplasia punctata, type 1, 215100 601769 Osteoporosis, involutional 601769 Rickets, vitamin D-resistant, 277440 601771 Glaucoma 3A, primary infantile, 231300 601780 Ceroid-lipofuscinosis, neuronal-6, variant late infantile 601785 Carbohydrate-deficient glycoprotein syndrome, type I, 212065 601843 Hypothyroidism, congenital, 274400 601846 Muscular dystrophy with rimmed vacuoles 601850 Retinitis pigmentosa-deafness syndrome 601863 Bare lymphocyte syndrome, complementation group C 601868 Deafness, autosomal dominant 13 601884 (High bone mass) 601885 Cataract, zonular pulverulent-2 601975 Ectodermal dysplasia/skin fragility syndrome 602025 Obesity/hyperinsulinism, susceptibility to 602026 Refsum disease, 266500 602078 Fibrosis of extraocular muscles, congenital, 2 60208 Nephronophthisis, infantile 602136 Refsum disease, 266510 602136 Refsum disease, infantile, 266510		
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601885 Cataract, zonular pulverulent-2 601975 Ectodermal dysplasia/skin fragility syndrome 602025 Obesity/hyperinsulinism, susceptibility to 602026 Refsum disease, 266500 602078 Fibrosis of extraocular muscles, congenital, 2 602088 Nephronophthisis, infantile 602094 Lipodystrophy, familial partial 602099 Amytrophic lateral sclerosis-5 602116 Glioma 602134 Tremor, familial essential, 2 602136 Refsum disease, infantile, 266510		Deafness, autosomal dominant 13
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602116 Glioma 602134 Tremor, familial essential, 2 602136 Refsum disease, infantile, 266510	602094	Lipodystrophy, familial partial
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Refsum disease, infantile, 266510	602116	
Refsum disease, infantile, 266510	602134	Tremor, familial essential, 2
	602136	
OUZ130 Zeilweger syndrome-1, 214100	602136	Zellweger syndrome-1, 214100
602136 Adrenoleukodystrophy, neonatal, 202370	602136	Adrenoleukodystrophy, neonatal, 202370
602216 Peutz-Jeghers syndrome, 175200	602216	

602221	Stem-cell leukemia/lymphoma syndrome
602225	Cone-rod retinal dystrophy-2, 120970
602225	Leber congenital amaurosis, type III
602229	Waardenburg-Shah syndrome, 277580
602235	Epilepsy, benign, neonatal, type 1, 121200
602280	Retinitis pigmentosa-14, 600132
602447	Coronary artery disease, susceptibility to
602475	Ossification of posterior longitudinal ligament of spine
602477	Febrile convulsions, familial, 2
602491	Hyperlipidemia, familial combined, 1
602544	Parkinson disease, juvenile, type 2, 600116
602631	Rhabdomyosarcoma, 268210
602631	Breast Cancer
602716	Nephrosis-1, congenital, Finnish type, 256300
602772	Retinitis pitmentosa-24
602782	Faisalabad histiocytosis
602783	Spastic paraplegia-7

Polynucleotide and Polypeptide Variants

The present invention is also directed to variants of the reproductive system associated polynucleotide sequence disclosed in SEQ ID NO:X or the complementary strand thereto, nucleotide sequences encoding the polypeptide of SEQ ID NO:Y, the nucleotide sequence of SEQ ID NO:X encoding the polypeptide sequence as defined in column 6 of Table 1A, nucleotide sequences encoding the polypeptide as defined in column 6 of Table 1A, the nucleotide sequence as defined in columns 8 and 9 of Table 2, nucleotide sequences encoding the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2, the nucleotide sequence as defined in column 6 of Table 1B, nucleotide sequences encoding the polypeptide encoded by the nucleotide sequence as defined in column 6 of Table 1B, nucleotide sequences encoding the polypeptide encoded by the nucleotide sequence as defined in column 6 of Table 1B, the cDNA sequence contained in Clone ID NO:Z, and/or nucleotide sequences encoding a polypeptide encoded by the cDNA sequence contained in Clone ID NO:Z.

[0114] The present invention also encompasses variants of the polypeptide sequence disclosed in SEQ ID NO:Y, a polypeptide sequence as defined in column 6 of Table 1A, a polypeptide sequence encoded by the polynucleotide sequence in SEQ

ID NO:X, a polypeptide sequence encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2, a polypeptide sequence encoded by the nucleotide sequence as defined in column 6 of Table 1B, a polypeptide sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X, and/or a polypeptide sequence encoded by the cDNA sequence contained in Clone ID NO:Z.

[0115] "Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

Thus, one aspect of the invention provides an isolated nucleic acid [0116] molecule comprising, or alternatively consisting of, a polynucleotide having a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence described in SEQ ID NO:X or contained in the cDNA sequence of Clone ID NO:Z; (b) a nucleotide sequence in SEQ ID NO:X or the cDNA in Clone ID NO:Z which encodes a mature reproductive system associated polypeptide; (c) a nucleotide sequence in SEQ ID NO:X or the cDNA sequence of Clone ID NO:Z, which encodes a biologically active fragment of a reproductive system associated polypeptide; (d) a nucleotide sequence in SEQ ID NO:X or the cDNA sequence of Clone ID NO:Z, which encodes an antigenic fragment of a reproductive system associated polypeptide; (e) a nucleotide sequence encoding a reproductive system associated polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; (f) a nucleotide sequence encoding a mature reproductive system associated polypeptide of the amino acid sequence of SEQ ID NO:Y or the amino acid sequence encoded by the cDNA in Clone ID NO:Z; (g) a nucleotide sequence encoding a biologically active fragment of a reproductive system associated polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; (h) a nucleotide sequence encoding an antigenic fragment of a reproductive system associated polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; and (i) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b), (c), (d), (e), (f), (g), or (h), above.

The present invention is also directed to nucleic acid molecules which [0117]comprise, or alternatively consist of, a nucleotide sequence which is at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100%, identical to, for example, any of the nucleotide sequences in (a), (b), (c), (d), (e), (f), (g), (h), or (i) above, the nucleotide coding sequence in SEQ ID NO:X or the complementary strand thereto, the nucleotide coding sequence of the cDNA contained in Clone ID NO:Z or the complementary strand thereto, a nucleotide sequence encoding the polypeptide of SEQ ID NO:Y, a nucleotide sequence encoding a polypeptide sequence encoded by the nucleotide sequence in SEQ ID NO:X, a polypeptide sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X, a nucleotide sequence encoding the polypeptide encoded by the cDNA contained in Clone ID NO:Z, the nucleotide coding sequence in SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto, a nucleotide sequence encoding the polypeptide encoded by the nucleotide sequence in SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto, the nucleotide coding sequence in SEQ ID NO:B as defined in column 6 of Table 1B or the complementary strand thereto, a nucleotide sequence encoding the polypeptide encoded by the nucleotide sequence in SEQ ID NO:B as defined in column 6 of Table 1B or the complementary strand thereto, the nucleotide sequence in SEQ ID NO:X encoding the polypeptide sequence as defined in column 6 of Table 1A or the complementary strand thereto, nucleotide sequences encoding a polypeptide as defined in column 6 of Table 1A or the complementary strand thereto, and/or polynucleotide fragments of any of these nucleic acid molecules (e.g., those fragments described herein). Polynucleotides which hybridize to the complement of these nucleic acid molecules under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention, as are polypeptides encoded by these polynucleotides and nucleic acids.

[0118] In a preferred embodiment, the invention encompasses nucleic acid molecules which comprise, or alternatively, consist of a polynucleotide which hybridizes under stringent hybridization conditions, or alternatively, under lower stringency conditions, to a polynucleotide in (a), (b), (c), (d), (e), (f), (g), (h), or (i) above, as are polypeptides encoded by these polynucleotides. In another preferred

embodiment, polynucleotides which hybridize to the complement of these nucleic acid molecules under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention, as are polypeptides encoded by these polynucleotides.

In another embodiment, the invention provides a purified protein comprising, or alternatively consisting of, a polypeptide having an amino acid sequence selected from the group consisting of: (a) the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; (b) the amino acid sequence of a mature reproductive system associated polypeptide having the amino acid sequence of SEQ ID NO:Y or the amino acid sequence encoded by the cDNA in Clone ID NO:Z; (c) the amino acid sequence of a biologically active fragment of a reproductive system associated polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence of an antigenic fragment of a reproductive system associated polypeptide having the complete amino acid sequence of an antigenic fragment of a reproductive system associated polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Y.

[0120] The present invention is also directed to proteins which comprise, or alternatively consist of, an amino acid sequence which is at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100%, identical to, for example, any of the amino acid sequences in (a), (b), (c), or (d), above, the amino acid sequence shown in SEQ ID NO:Y, the amino acid sequence encoded by the cDNA contained in Clone ID NO:Z, the amino acid sequence of the polypeptide encoded by the nucleotide sequence in SEQ ID NO:X as defined in columns 8 and 9 of Table 2, the amino acid sequence of the polypeptide encoded by the nucleotide sequence in SEQ ID NO:B as defined in column 6 of Table 1B, the amino acid sequence as defined in column 6 of Table 1A, an amino acid sequence encoded by the nucleotide sequence in SEQ ID NO:X, and an amino acid sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X. Fragments of these polypeptides are also provided (e.g., those fragments described herein). Further proteins encoded by polynucleotides which hybridize to the complement of the nucleic acid molecules encoding these amino acid sequences under stringent hybridization conditions or alternatively, under lower

stringency conditions, are also encompassed by the invention, as are the polynucleotides encoding these proteins.

"identical" to a reference nucleotide sequence of the present invention, it is intended that the nucleotide sequence of the nucleic acid is identical to the reference sequence except that the nucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the polypeptide. In other words, to obtain a nucleic acid having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may be inserted into the reference sequence. The query sequence may be an entire sequence referred to in Table 1A or 2 as the ORF (open reading frame), or any fragment specified, as described herein.

As a practical matter, whether any particular nucleic acid molecule or [0122] polypeptide is at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleotide sequence of the present invention can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci. 6:237-245 (1990)). In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by converting U's to T's. The result of said global sequence alignment is expressed as percent identity. Preferred parameters used in a FASTDB alignment of DNA sequences to calculate percent identity are: Matrix=Unitary, k-tuple=4, Mismatch Penalty=1, Joining Penalty=30, Randomization Group Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty 0.05, Window Size=500 or the length of the subject nucleotide sequence, whichever is shorter.

[0123] If the subject sequence is shorter than the query sequence because of 5' or 3' deletions, not because of internal deletions, a manual correction must be made to the results. This is because the FASTDB program does not account for 5' and 3'

truncations of the subject sequence when calculating percent identity. For subject sequences truncated at the 5' or 3' ends, relative to the query sequence, the percent identity is corrected by calculating the number of bases of the query sequence that are 5' and 3' of the subject sequence, which are not matched/aligned, as a percent of the total bases of the query sequence. Whether a nucleotide is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This corrected score is what is used for the purposes of the present invention. Only bases outside the 5' and 3' bases of the subject sequence, as displayed by the FASTDB alignment, which are not matched/aligned with the query sequence, are calculated for the purposes of manually adjusting the percent identity score.

[0124] For example, a 90 base subject sequence is aligned to a 100 base query sequence to determine percent identity. The deletions occur at the 5' end of the subject sequence and therefore, the FASTDB alignment does not show a matched/alignment of the first 10 bases at 5' end. The 10 unpaired bases represent 10% of the sequence (number of bases at the 5' and 3' ends not matched/total number of bases in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 bases were perfectly matched the final percent identity would be 90%. In another example, a 90 base subject sequence is compared with a 100 base query sequence. This time the deletions are internal deletions so that there are no bases on the 5' or 3' of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only bases 5' and 3' of the subject sequence which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to be made for the purposes of the present invention.

[0125] By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a query amino acid sequence of the present invention, it is intended that the amino acid sequence of the subject polypeptide is identical to the query sequence except that the subject polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the query amino acid sequence. In other

words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a query amino acid sequence, up to 5% of the amino acid residues in the subject sequence may be inserted, deleted, (indels) or substituted with another amino acid. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

[0126] As a practical matter, whether any particular polypeptide is at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to, for instance, the amino acid sequence of a polypeptide referred to in Table 1A (e.g., an amino acid sequence identified in columns 5 or 6) or Table 2 (e.g., the amino acid sequence of the polypeptide encoded by the polynucleotide sequence defined in columns 8 and 9 of Table 2) or a fragment thereof, the amino acid sequence of the polypeptide encoded by the polynucleotide sequence in SEQ ID NO:B as defined in column 6 of Table 1B or a fragment thereof, the amino acid sequence of the polypeptide encoded by the nucleotide sequence in SEQ ID NO:X or a fragment thereof, or an amino acid sequence of the polypeptide encoded by cDNA contained in Clone ID NO:Z, or a fragment thereof, can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci.6:237-245 (1990)). In a sequence alignment the query and subject sequences are either both nucleotide sequences or both amino acid sequences. The result of said global sequence alignment is expressed as percent identity. Preferred parameters used in a FASTDB amino acid alignment are: Matrix=PAM 0, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group Length=0, Cutoff Score=1, Window Size=sequence length, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the length of the subject amino acid sequence, whichever is shorter.

[0127] If the subject sequence is shorter than the query sequence due to N- or C-terminal deletions, not because of internal deletions, a manual correction must be made to the results. This is because the FASTDB program does not account for N-

and C-terminal truncations of the subject sequence when calculating global percent identity. For subject sequences truncated at the N- and C-termini, relative to the query sequence, the percent identity is corrected by calculating the number of residues of the query sequence that are N- and C-terminal of the subject sequence, which are not matched/aligned with a corresponding subject residue, as a percent of the total bases of the query sequence. Whether a residue is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This final percent identity score is what is used for the purposes of the present invention. Only residues to the N- and C-termini of the subject sequence, which are not matched/aligned with the query sequence, are considered for the purposes of manually adjusting the percent identity score. That is, only query residue positions outside the farthest N- and C- terminal residues of the subject sequence.

[0128] For example, a 90 amino acid residue subject sequence is aligned with a 100 residue query sequence to determine percent identity. The deletion occurs at the N-terminus of the subject sequence and therefore, the FASTDB alignment does not show a matching/alignment of the first 10 residues at the N-terminus. The 10 unpaired residues represent 10% of the sequence (number of residues at the N- and C- termini not matched/total number of residues in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 residues were perfectly matched the final percent identity would be 90%. In another example, a 90 residue subject sequence is compared with a 100 residue query sequence. This time the deletions are internal deletions so there are no residues at the N- or C-termini of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only residue positions outside the N- and C-terminal ends of the subject sequence, as displayed in the FASTDB alignment, which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to be made for the purposes of the present invention.

[0129] The polynucleotide variants of the invention may contain alterations in the coding regions, non-coding regions, or both. Especially preferred are polynucleotide

variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, polypeptide variants in which less than 50, less than 40, less than 30, less than 20, less than 10, or 5-50, 5-25, 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in the human mRNA to those preferred by a bacterial host such as *E. coli*).

- [0130] Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level and are included in the present invention. Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.
- technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the polypeptides of the present invention without substantial loss of biological function. As an example, the authors of Ron et al., J. Biol. Chem. 268: 2984-2988 (1993), reported variant KGF proteins having heparin binding activity even after deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).)
- [0132] Moreover, ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (J. Biol. Chem. 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1a. They used random mutagenesis to generate over 3,500 individual IL-1a mutants that averaged 2.5 amino acid changes per variant over the entire length of the molecule. Multiple mutations were examined at every possible amino acid position. The investigators found that "[m]ost of the

molecule could be altered with little effect on either [binding or biological activity]." In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained. For example, the ability of a deletion variant to induce and/or to bind antibodies which recognize the secreted form will likely be retained when less than the majority of the residues of the secreted form are removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by routine methods described herein and otherwise known in the art.

[0134] Thus, the invention further includes polypeptide variants which show a functional activity (e.g., biological activity) of the polypeptides of the invention. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity.

[0135] The present application is directed to nucleic acid molecules at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100% identical to the nucleic acid sequences disclosed herein, (e.g., encoding a polypeptide having the amino acid sequence of an N and/or C terminal deletion), irrespective of whether they encode a polypeptide having functional activity. This is because even where a particular nucleic acid molecule does not encode a polypeptide having functional activity, one of skill in the art would still know how to use the nucleic acid molecule, for instance, as a hybridization probe or a polymerase chain reaction (PCR) primer. Uses of the nucleic acid molecules of the present invention that do not encode a polypeptide having functional activity include, inter alia, (1) isolating a gene or allelic or splice variants thereof in a cDNA library; (2) in situ hybridization (e.g., "FISH") to metaphase chromosomal spreads to provide precise chromosomal location of the gene, as described in Verma et al., Human Chromosomes: A Manual of Basic Techniques, Pergamon Press, New York (1988); (3) Northern Blot analysis for detecting mRNA expression in specific tissues (e.g., normal reproductive system tissues or diseased

reproductive system tissues); and (4) in situ hybridization (e.g., histochemistry) for detecting mRNA expression in specific tissues (e.g., normal reproductive system tissues or diseased reproductive system tissues).

[0136] Preferred, however, are nucleic acid molecules having sequences at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100% identical to the nucleic acid sequences disclosed herein, which do, in fact, encode a polypeptide having functional activity. By a polypeptide having "functional activity" is meant, a polypeptide capable of displaying one or more known functional activities associated with a full-length (complete) protein of the invention. Such functional activities include, but are not limited to, biological activity, antigenicity [ability to bind (or compete with a polypeptide of the invention for binding) to an anti-polypeptide of the invention antibody], immunogenicity (ability to generate antibody which binds to a specific polypeptide of the invention), ability to form multimers with polypeptides of the invention.

[0137] The functional activity of the polypeptides, and fragments, variants and derivatives of the invention, can be assayed by various methods.

For example, in one embodiment where one is assaying for the ability to [0138] bind or compete with full-length polypeptide of the present invention for binding to an anti-polypeptide of the invention antibody, various immunoassays known in the art can be used, including but not limited to, competitive and non-competitive assay systems using techniques such as radioimmunoassays, ELISA (enzyme linked immunosorbent assay), "sandwich" immunoassays, immunoradiometric assays, gel diffusion precipitation reactions, immunodiffusion assays, in situ immunoassays (using colloidal gold, enzyme or radioisotope labels, for example), western blots, precipitation reactions, agglutination assays (e.g., gel agglutination assays, hemagglutination assays), complement fixation assays, immunofluorescence assays, protein A assays, and immunoelectrophoresis assays, etc. In one embodiment, antibody binding is detected by detecting a label on the primary antibody. In another embodiment, the primary antibody is detected by detecting binding of a secondary antibody or reagent to the primary antibody. In a further embodiment, the secondary antibody is labeled. Many means are known in the art for detecting binding in an immunoassay and are within the scope of the present invention.

[0139] In another embodiment, where a ligand is identified, or the ability of a polypeptide fragment, variant or derivative of the invention to multimerize is being evaluated, binding can be assayed, e.g., by means well-known in the art, such as, for example, reducing and non-reducing gel chromatography, protein affinity chromatography, and affinity blotting. See generally, Phizicky et al., Microbiol. Rev. 59:94-123 (1995). In another embodiment, the ability of physiological correlates of a polypeptide of the present invention to bind to a substrate(s) of the polypeptide of the invention can be routinely assayed using techniques known in the art.

- [0140] In addition, assays described herein (see Examples) and otherwise known in the art may routinely be applied to measure the ability of polypeptides of the present invention and fragments, variants and derivatives thereof to elicit polypeptide related biological activity (either *in vitro* or *in vivo*). Other methods will be known to the skilled artisan and are within the scope of the invention.
- Of course, due to the degeneracy of the genetic code, one of ordinary skill [0141] in the art will immediately recognize that a large number of the nucleic acid molecules having a sequence at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99%, or 100% identical to, for example, the nucleic acid sequence of the cDNA contained in Clone ID NO:Z, a nucleic acid sequence referred to in Table 1A (e.g., SEQ ID NO:X), a nucleic acid sequence disclosed in Table 2 (e.g., the nucleic acid sequence delineated in columns 8 and 9) or fragments thereof, will encode polypeptides "having functional activity." In fact, since degenerate variants of any of these nucleotide sequences all encode the same polypeptide, in many instances, this will be clear to the skilled artisan even without performing the above described comparison assay. It will be further recognized in the art that, for such nucleic acid molecules that are not degenerate variants, a reasonable number will also encode a polypeptide having functional activity. This is because the skilled artisan is fully aware of amino acid substitutions that are either less likely or not likely to significantly effect protein function (e.g., replacing one aliphatic amino acid with a second aliphatic amino acid), as further described below.
- [0142] For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie et al., "Deciphering the Message in Protein Sequences: Tolerance to Amino Acid Substitutions," Science 247:1306-1310

(1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

[0143] The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

[0144] The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. See Cunningham et al., Science 244:1081-1085 (1989). The resulting mutant molecules can then be tested for biological activity.

[0145] As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly. Besides conservative amino acid substitutions, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitutions with one or more of the amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example,

polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as, for example, an IgG Fc fusion region peptide, serum albumin (preferably human serum albumin) or a fragment or variant thereof, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

[0146] For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. See Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).

A further embodiment of the invention relates to polypeptides which comprise the amino acid sequence of a polypeptide having an amino acid sequence which contains at least one amino acid substitution, but not more than 50 amino acid substitutions, even more preferably, not more than 40 amino acid substitutions, still more preferably, not more than 30 amino acid substitutions, and still even more preferably, not more than 20 amino acid substitutions from a polypeptide sequence disclosed herein. Of course it is highly preferable for a polypeptide to have an amino acid sequence which comprises the amino acid sequence of a polypeptide of SEQ ID NO:Y, an amino acid sequence encoded by SEQ ID NO:X, an amino acid sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, an amino acid sequence encoded by the complement of SEQ ID NO:X, and/or the amino acid sequence encoded by cDNA contained in Clone ID NO:Z which contains, in order of ever-increasing preference, at least one, but not more than 10, 9, 8, 7, 6, 5, 4, 3, 2 or 1 amino acid substitutions.

In specific embodiments, the polypeptides of the invention comprise, or alternatively, consist of, fragments or variants of a reference amino acid sequence selected from: (a) the amino acid sequence of SEQ ID NO:Y or fragments thereof (e.g., the mature form and/or other fragments described herein); (b) the amino acid sequence encoded by SEQ ID NO:X or fragments thereof; (c) the amino acid sequence encoded by the complement of SEQ ID NO:X or fragments thereof; (d) the amino acid

sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or fragments thereof; and (e) the amino acid sequence encoded by cDNA contained in Clone ID NO:Z or fragments thereof; wherein the fragments or variants have 1-5, 5-10, 5-25, 5-50, 10-50 or 50-150, amino acid residue additions, substitutions, and/or deletions when compared to the reference amino acid sequence. In preferred embodiments, the amino acid substitutions are conservative. Polynucleotides encoding these polypeptides are also encompassed by the invention.

Polynucleotide and Polypeptide Fragments

The present invention is also directed to polynucleotide fragments of the polynucleotides (nucleic acids) of the invention. In the present invention, a "polynucleotide fragment" refers to a polynucleotide having a nucleic acid sequence which, for example: is a portion of the cDNA contained in Clone ID NO:Z or the complementary strand thereto; is a portion of the polynucleotide sequence encoding the polypeptide encoded by the cDNA contained in Clone ID NO:Z or the complementary strand thereto; is a portion of a polynucleotide sequence encoding the amino acid sequence encoded by the region of SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto; is a portion of the polynucleotide sequence of SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto; is a portion of the polynucleotide sequence in SEQ ID NO:X or the complementary strand thereto; is a polynucleotide sequence encoding a portion of the polypeptide of SEQ ID NO:Y; is a polynucleotide sequence encoding a portion of a polypeptide encoded by SEQ ID NO:X; is a polynucleotide sequence encoding a portion of a polypeptide encoded by the complement of the polynucleotide sequence in SEQ ID NO:X; is a portion of a polynucleotide sequence encoding the amino acid sequence encoded by the region of SEQ ID NO:B as defined in column 6 of Table 1B or the complementary strand thereto; or is a portion of the polynucleotide sequence of SEQ ID NO:B as defined in column 6 of Table 1B or the complementary strand thereto.

[0150] The polynucleotide fragments of the invention are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt, at least about 50 nt, at least about 75

nt, or at least about 150 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in Clone ID NO:Z, or the nucleotide sequence shown in SEQ ID NO:X or the complementary stand thereto. In this context "about" includes the particularly recited value or a value larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. These nucleotide fragments have uses that include, but are not limited to, as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., at least 160, 170, 180, 190, 200, 250, 500, 600, 1000, or 2000 nucleotides in length) are also encompassed by the invention.

Moreover, representative examples of polynucleotide fragments of the [0151] invention, comprise, or alternatively consist of, a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, 701-750, 751-800, 800-850, 851-900, 901-950, 951-1000, 1001-1050, 1051-1100, 1101-1150, 1151-1200, 1201-1250, 1251-1300, 1301-1350, 1351-1400, 1401-1450, 1451-1500, 1501-1550, 1551-1600, 1601-1650, 1651-1700, 1701-1750, 1751-1800, 1801-1850, 1851-1900, 1901-1950, 1951-2000, 2001-2050, 2051-2100, 2101-2150, 2151-2200, 2201-2250, 2251-2300, 2301-2350, 2351-2400, 2401-2450, 2451-2500, 2501-2550, 2551-2600, 2601-2650, 2651-2700, 2701-2750, 2751-2800, 2801-2850, 2851-2900, 2901-2950, 2951-3000, 3001-3050, 3051-3100, 3101-3150, 3151-3200, 3201-3250, 3251-3300, 3301-3350, 3351-3400, 3401-3450, 3451-3500, 3501-3550, 3551-3600, 3601-3650, 3651-3700, 3701-3750, 3751-3800, 3801-3850, 3851-3900, 3901-3950, 3951-4000, 4001-4050, 4051-4100, 4101-4150, 4151-4200, 4201-4250, 4251-4300, 4301-4350, 4351-4400, 4401-4450, 4451-4500, 4501-4550, 4551-4600, 4601-4650, 4651-4700, 4701-4750, 4751-4800, 4801-4850, 4851-4900, 4901-4950, 4951-5000, 5001-5050, 5051-5100, 5101-5150, 5151-5200, 5201-5250, 5251-5300, 5301-5350, 5351-5400, 5401-5450, 5451-5500, 5501-5550, 5551-5600, 5601-5650, 5651-5700, 5701-5750, 5751-5800, 5801-5850, 5851-5900, 5901-5950, 5951-6000, 6001-6050, 6051-6100, 6101-6150, 6151-6200, 6201-6250, 6251-6300, 6301-6350, 6351-6400, 6401-6450, 6451-6500, 6501-6550, 6551-6600, 6601-6650, 6651-6700, 6701-6750, 6751-6800, 6801-6850, 6851-6900, 6901-6950, 6951-7000, 7001-7050, 7051-7100, 7101-7150, 7151-7200, 7201-7250, 7251-7300 or 7301 to the end of SEQ ID NO:X, or the complementary strand

thereto. In this context "about" includes the particularly recited range or a range larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has a functional activity (e.g., biological activity). More preferably, these polynucleotides can be used as probes or primers as discussed herein. Polynucleotides which hybridize to one or more of these polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions are also encompassed by the invention, as are polypeptides encoded by these polynucleotides.

[0152] Further representative examples of polynucleotide fragments of the invention, comprise, or alternatively consist of, a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, 701-750, 751-800, 800-850, 851-900, 901-950, 951-1000, 1001-1050, 1051-1100, 1101-1150, 1151-1200, 1201-1250, 1251-1300, 1301-1350, 1351-1400, 1401-1450, 1451-1500, 1501-1550, 1551-1600, 1601-1650, 1651-1700, 1701-1750, 1751-1800, 1801-1850, 1851-1900, 1901-1950, 1951-2000, 2001-2050, 2051-2100, 2101-2150, 2151-2200, 2201-2250, 2251-2300, 2301-2350, 2351-2400, 2401-2450, 2451-2500, 2501-2550, 2551-2600, 2601-2650, 2651-2700, 2701-2750, 2751-2800, 2801-2850, 2851-2900, 2901-2950, 2951-3000, 3001-3050, 3051-3100, 3101-3150, 3151-3200, 3201-3250, 3251-3300, 3301-3350, 3351-3400, 3401-3450, 3451-3500, 3501-3550, 3551-3600, 3601-3650, 3651-3700, 3701-3750, 3751-3800, 3801-3850, 3851-3900, 3901-3950, 3951-4000, 4001-4050, 4051-4100, 4101-4150, 4151-4200, 4201-4250, 4251-4300, 4301-4350, 4351-4400, 4401-4450, 4451-4500, 4501-4550, 4551-4600, 4601-4650, 4651-4700, 4701-4750, 4751-4800, 4801-4850, 4851-4900, 4901-4950, 4951-5000, 5001-5050, 5051-5100, 5101-5150, 5151-5200, 5201-5250, 5251-5300, 5301-5350, 5351-5400, 5401-5450, 5451-5500, 5501-5550, 5551-5600, 5601-5650, 5651-5700, 5701-5750, 5751-5800, 5801-5850, 5851-5900, 5901-5950, 5951-6000, 6001-6050, 6051-6100, 6101-6150, 6151-6200, 6201-6250, 6251-6300, 6301-6350, 6351-6400, 6401-6450, 6451-6500, 6501-6550, 6551-6600, 6601-6650, 6651-6700, 6701-6750, 6751-6800, 6801-6850, 6851-6900, 6901-6950, 6951-7000, 7001-7050, 7051-7100, 7101-7150, 7151-7200, 7201-7250, 7251-7300 or 7301 to the end of the cDNA sequence contained in Clone ID NO:Z, or the complementary strand thereto. In this context "about" includes the

particularly recited range or a range larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has a functional activity (e.g., biological activity). More preferably, these polynucleotides can be used as probes or primers as discussed herein. Polynucleotides which hybridize to one or more of these polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions are also encompassed by the invention, as are polypeptides encoded by these polynucleotides.

Moreover, representative examples of polynucleotide fragments of the [0153] invention comprise, or alternatively consist of, a nucleic acid sequence comprising one, two, three, four, five, six, seven, eight, nine, ten, or more of the above described polynucleotide fragments of the invention in combination with a polynucleotide sequence delineated in Table 1B column 6. Additional, representative examples of polynucleotide fragments of the invention comprise, or alternatively consist of, a nucleic acid sequence comprising one, two, three, four, five, six, seven, eight, nine, ten, or more of the above described polynucleotide fragments of the invention in combination with a polynucleotide sequence that is the complementary strand of a sequence delineated in column 6 of Table 1B. In further embodiments, the abovedescribed polynucleotide fragments of the invention comprise, or alternatively consist of, sequences delineated in Table 1B, column 6, and have a nucleic acid sequence which is different from that of the BAC fragment having the sequence disclosed in SEQ ID NO:B (see Table 1B, column 5). In additional embodiments, the abovedescribed polynucleotide fragments of the invention comprise, or alternatively consist of, sequences delineated in Table 1B, column 6, and have a nucleic acid sequence which is different from that published for the BAC clone identified as BAC ID NO:A (see Table 1B, column 4). In additional embodiments, the above-described polynucleotides of the invention comprise, or alternatively consist of, sequences delineated Table 1B, column 6, and have a nucleic acid sequence which is different from that contained in the BAC clone identified as BAC ID NO:A (see Table 1B, column 4). Polypeptides encoded by these polynucleotides, other polynucleotides that encode these polypeptides, and antibodies that bind these polypeptides are also

encompassed by the invention. Additionally, fragments and variants of the above-described polynucleotides and polypeptides are also encompassed by the invention.

[0154] In additional specific embodiments, polynucleotides of the invention comprise, or alternatively consist of, one, two, three, four, five, six, seven, eight, nine, ten, or more fragments of the sequences delineated in column 6 of Table 1B, and the polynucleotide sequence of SEQ ID NO:X (e.g., as defined in Table 1B, column 2) or fragments or variants thereof. Polypeptides encoded by these polynucleotides, other polynucleotides that encode these polypeptides, and antibodies that bind these polypeptides are also encompassed by the invention.

[0155] In additional specific embodiments, polynucleotides of the invention comprise, or alternatively consist of, one, two, three, four, five, six, seven, eight, nine, ten, or more fragments of the sequences delineated in column 6 of Table 1B which correspond to the same Clone ID NO:Z (see Table 1B, column 1), and the polynucleotide sequence of SEQ ID NO:X (e.g., as defined in Table 1A or 1B) or fragments or variants thereof. Polypeptides encoded by these polynucleotides, other polynucleotides that encode these polypeptides, and antibodies that bind these polypeptides are also encompassed by the invention.

[0156] In further specific embodiments, polynucleotides of the invention comprise, or alternatively consist of, one, two, three, four, five, six, seven, eight, nine, ten, or more fragments of the sequences delineated in the same row of column 6 of Table 1B, and the polynucleotide sequence of SEQ ID NO:X (e.g., as defined in Table 1A or 1B) or fragments or variants thereof. Polypeptides encoded by these polynucleotides, other polynucleotides that encode these polypeptides, and antibodies that bind these polypeptides are also encompassed by the invention.

[0157] In additional specific embodiments, polynucleotides of the invention comprise, or alternatively consist of a polynucleotide sequence in which the 3' 10 polynucleotides of one of the sequences delineated in column 6 of Table 1B and the 5' 10 polynucleotides of the sequence of SEQ ID NO:X are directly contiguous. Nucleic acids which hybridize to the complement of these 20 contiguous polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention. Polypeptides encoded by these polynucleotides and/or nucleic acids, other polynucleotides and/or nucleic acids that

encode these polypeptides, and antibodies that bind these polypeptides are also encompassed by the invention. Additionally, fragments and variants of the above-described polynucleotides, nucleic acids, and polypeptides are also encompassed by the invention.

[0158] In additional specific embodiments, polynucleotides of the invention comprise, or alternatively consist of a polynucleotide sequence in which the 3' 10 polynucleotides of one of the sequences delineated in column 6 of Table 1B and the 5' 10 polynucleotides of a fragment or variant of the sequence of SEQ ID NO:X (e.g., as described herein) are directly contiguous Nucleic acids which hybridize to the complement of these 20 contiguous polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention. Polypeptides encoded by these polynucleotides and/or nucleic acids, other polynucleotides and/or nucleic acids encoding these polypeptides, and antibodies that bind these polypeptides are also encompassed by the invention. Additionally, fragments and variants of the above-described polynucleotides, nucleic acids, and polypeptides are also encompassed by the invention.

[0159] In further specific embodiments, polynucleotides of the invention comprise, or alternatively consist of a polynucleotide sequence in which the 3' 10 polynucleotides of a fragment or variant of the sequence of SEQ ID NO:X and the 5' 10 polynucleotides of the sequence of one of the sequences delineated in column 6 of Table 1B are directly contiguous. Nucleic acids which hybridize to the complement of these 20 contiguous polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention. Polypeptides encoded by these polynucleotides and/or nucleic acids, other polynucleotides and/or nucleic acids encoding these polypeptides, and antibodies that bind these polypeptides are also encompassed by the invention. Additionally, fragments and variants of the above-described polynucleotides, nucleic acids, and polypeptides are also encompassed by the invention.

[0160] In specific embodiments, polynucleotides of the invention comprise, or alternatively consist of a polynucleotide sequence in which the 3' 10 polynucleotides of one of the sequences delineated in column 6 of Table 1B and the 5' 10 polynucleotides of another sequence in column 6 are directly contiguous. In preferred

embodiments, the 3' 10 polynucleotides of one of the sequences delineated in column 6 of Table 1B is directly contiguous with the 5' 10 polynucleotides of the next sequential exon delineated in Table 1B, column 6. Nucleic acids which hybridize to the complement of these 20 contiguous polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention. Polypeptides encoded by these polynucleotides and/or nucleic acids, other polynucleotides and/or nucleic acids encoding these polypeptides, and antibodies that bind these polypeptides are also encompassed by the invention. Additionally, fragments and variants of the above-described polynucleotides, nucleic acids, and polypeptides are also encompassed by the invention.

In the present invention, a "polypeptide fragment" refers to an amino acid sequence which is a portion of that contained in SEQ ID NO:Y, a portion of an amino acid sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, a portion of an amino acid sequence encoded by the polynucleotide sequence of SEQ ID NO:X, a portion of an amino acid sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X, and/or a portion of an amino acid sequence encoded by the cDNA contained in Clone ID NO:Z. Protein (polypeptide) fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single Representative examples of polypeptide fragments of the continuous region. invention, include, for example, fragments comprising, or alternatively consisting of, from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 102-120, 121-140, 141-160, 161-180, 181-200, 201-220, 221-240, 241-260, 261-280, 281-300, 301-320, 321-340, 341-360, 361-380, 381-400, 401-420, 421-440, 441-460, 461-480, 481-500, 501-520, 521-540, 541-560, 561-580, 581-600, 601-620, 621-640, 641-660, 661-680, 681-700, 701-720, 721-740, 741-760, 761-780, 781-800, 801-820, 821-840, 841-860, 861-880, 881-900, 901-920, 921-940, 941-960, 961-980, 981-1000, 1001-1020, 1021-1040, 1041-1060, 1061-1080, 1081-1100, 1101-1120, 1121-1140, 1141-1160, 1161-1180, 1181-1200, 1201-1220, 1221-1240, 1241-1260, 1261-1280, 1281-1300, 1301-1320, 1321-1340, 1341-1360, 1361-1380, 1381-1400, 1401-1420, 1421-1440, or 1441 to the end of the coding region. In a preferred embodiment, polypeptide fragments of the invention include, for example, fragments comprising, or alternatively consisting

of, from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 102-120, 121-140, 141-160, 161-180, 181-200, 201-220, 221-240, 241-260, 261-280, 281-300, 301-320, 321-340, 341-360, 361-380, 381-400, 401-420, 421-440, 441-460, 461-480, 481-500, 501-520, 521-540, 541-560, 561-580, 581-600, 601-620, 621-640, 641-660, 661-680, 681-700, 701-720, 721-740, 741-760, 761-780, 781-800, 801-820, 821-840, 841-860, 861-880, 881-900, 901-920, 921-940, 941-960, 961-980, 981-1000, 1001-1020, 1021-1040, 1041-1060, 1061-1080, 1081-1100, 1101-1120, 1121-1140, 1141-1160, 1161-1180, 1181-1200, 1201-1220, 1221-1240, 1241-1260, 1261-1280, 1281-1300, 1301-1320, 1321-1340, 1341-1360, 1361-1380, 1381-1400, 1401-1420, 1421-1440, or 1441 to the end of the coding region of SEQ ID NO:Y. Moreover, polypeptide fragments of the invention may be at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about" includes the particularly recited ranges or values, or ranges or values larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes. Polynucleotides encoding these polypeptide fragments are also encompassed by the invention.

Even if deletion of one or more amino acids from the N-terminus of a protein results in modification of loss of one or more biological functions of the protein, other functional activities (e.g., biological activities, ability to multimerize, ability to bind a ligand) may still be retained. For example, the ability of shortened muteins to induce and/or bind to antibodies which recognize the complete or mature forms of the polypeptides generally will be retained when less than the majority of the residues of the complete or mature polypeptide are removed from the N-terminus. Whether a particular polypeptide lacking N-terminal residues of a complete polypeptide retains such immunologic activities can readily be determined by routine methods described herein and otherwise known in the art. It is not unlikely that a mutein with a large number of deleted N-terminal amino acid residues may retain some biological or immunogenic activities. In fact, peptides composed of as few as six amino acid residues may often evoke an immune response.

[0163] Accordingly, polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or

the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions is preferred. Similarly, polynucleotides encoding these polypeptide fragments are also preferred.

The present invention further provides polypeptides having one or more residues deleted from the amino terminus of the amino acid sequence of a polypeptide disclosed herein (e.g., a polypeptide of SEQ ID NO:Y, a polypeptide encoded by the polynucleotide sequence contained in SEQ ID NO:X or the complement thereof, a polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, a polypeptide encoded by the portion of SEQ ID NO:B as defined in column 6 of Table 1B, and/or a polypeptide encoded by the cDNA contained in Clone ID NO:Z). In particular, N-terminal deletions may be described by the general formula m-q, where q is a whole integer representing the total number of amino acid residues in a polypeptide of the invention (e.g., the polypeptide disclosed in SEQ ID NO:Y, or the polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2), and m is defined as any integer ranging from 2 to q-6. Polynucleotides encoding these polypeptides are also encompassed by the invention.

The present invention further provides polypeptides having one or more residues from the carboxy terminus of the amino acid sequence of a polypeptide disclosed herein (e.g., a polypeptide of SEQ ID NO:Y, a polypeptide encoded by the polynucleotide sequence contained in SEQ ID NO:X, a polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or a polypeptide encoded by the cDNA contained in Clone ID NO:Z). In particular, C-terminal deletions may be described by the general formula 1-n, where n is any whole integer ranging from 6 to q-1, and where n corresponds to the position of amino acid residue in a polypeptide of the invention. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0166] In addition, any of the above described N- or C-terminal deletions can be combined to produce a N- and C-terminal deleted polypeptide. The invention also

provides polypeptides having one or more amino acids deleted from both the amino and the carboxyl termini, which may be described generally as having residues m-n of a polypeptide encoded by SEQ ID NO:X (e.g., including, but not limited to, the preferred polypeptide disclosed as SEQ ID NO:Y and the polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2), the cDNA contained in Clone ID NO:Z, and/or the complement thereof, where n and m are integers as described above. Polynucleotides encoding these polypeptides are also encompassed by the invention.

the C-terminus of a protein results in modification of loss of one or more biological functions of the protein, other functional activities (e.g., biological activities, ability to multimerize, ability to bind a ligand) may still be retained. For example the ability of the shortened mutein to induce and/or bind to antibodies which recognize the complete or mature forms of the polypeptide generally will be retained when less than the majority of the residues of the complete or mature polypeptide are removed from the C-terminus. Whether a particular polypeptide lacking C-terminal residues of a complete polypeptide retains such immunologic activities can readily be determined by routine methods described herein and otherwise known in the art. It is not unlikely that a mutein with a large number of deleted C-terminal amino acid residues may retain some biological or immunogenic activities. In fact, peptides composed of as few as six amino acid residues may often evoke an immune response.

The present application is also directed to proteins containing polypeptides at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to a polypeptide sequence set forth herein. In preferred embodiments, the application is directed to proteins containing polypeptides at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to polypeptides having the amino acid sequence of the specific N- and C-terminal deletions. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0169] Any polypeptide sequence encoded by, for example, the polynucleotide sequences set forth as SEQ ID NO:X or the complement thereof, (presented, for example, in Tables 1A and 2), the cDNA contained in Clone ID NO:Z, or the polynucleotide sequence as defined in column 6 of Table 1B, may be analyzed to

determine certain preferred regions of the polypeptide. For example, the amino acid sequence of a polypeptide encoded by a polynucleotide sequence of SEQ ID NO:X (e.g., the polypeptide of SEQ ID NO:Y and the polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2) or the cDNA contained in Clone ID NO:Z may be analyzed using the default parameters of the DNASTAR computer algorithm (DNASTAR, Inc., 1228 S. Park St., Madison, WI 53715 USA; http://www.dnastar.com/).

[0170] Polypeptide regions that may be routinely obtained using the DNASTAR computer algorithm include, but are not limited to, Garnier-Robson alpha-regions, beta-regions, turn-regions, and coil-regions; Chou-Fasman alpha-regions, beta-regions, and turn-regions; Kyte-Doolittle hydrophilic regions and hydrophobic regions; Eisenberg alpha- and beta-amphipathic regions; Karplus-Schulz flexible regions; Emini surface-forming regions; and Jameson-Wolf regions of high antigenic index. Among highly preferred polynucleotides of the invention in this regard are those that encode polypeptides comprising regions that combine several structural features, such as several (e.g., 1, 2, 3 or 4) of the features set out above.

[0171] Additionally, Kyte-Doolittle hydrophilic regions and hydrophobic regions, Emini surface-forming regions, and Jameson-Wolf regions of high antigenic index (i.e., containing four or more contiguous amino acids having an antigenic index of greater than or equal to 1.5, as identified using the default parameters of the Jameson-Wolf program) can routinely be used to determine polypeptide regions that exhibit a high degree of potential for antigenicity. Regions of high antigenicity are determined from data by DNASTAR analysis by choosing values which represent regions of the polypeptide which are likely to be exposed on the surface of the polypeptide in an environment in which antigen recognition may occur in the process of initiation of an immune response.

[0172] Preferred polypeptide fragments of the invention are fragments comprising, or alternatively, consisting of, an amino acid sequence that displays a functional activity (e.g. biological activity) of the polypeptide sequence of which the amino acid sequence is a fragment. By a polypeptide displaying a "functional activity" is meant a polypeptide capable of one or more known functional activities associated with a full-

length protein, such as, for example, biological activity, antigenicity, immunogenicity, and/or multimerization, as described herein.

[0173] Other preferred polypeptide fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

[0174] In preferred embodiments, polypeptides of the invention comprise, or alternatively consist of, one, two, three, four, five or more of the antigenic fragments of the polypeptide of SEQ ID NO:Y, or portions thereof. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0175] The present invention encompasses polypeptides comprising, or alternatively consisting of, an epitope of: the polypeptide sequence shown in SEQ ID NO:Y; a polypeptide sequence encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2; the polypeptide sequence encoded by the portion of SEQ ID NO:B as defined in column 6 of Table 1B or the complement thereto; the polypeptide sequence encoded by the cDNA contained in Clone ID NO:Z; or the polypeptide sequence encoded by a polynucleotide that hybridizes to the sequence of SEQ ID NO:X, the complement of the sequence of SEQ ID NO:X, the complement of a portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, or the cDNA sequence contained in Clone ID NO:Z under stringent hybridization conditions or alternatively, under lower stringency hybridization as defined supra. The present invention further encompasses polynucleotide sequences encoding an epitope of a polypeptide sequence of the invention (such as, for example, the sequence disclosed in SEQ ID NO:X, or a fragment thereof), polynucleotide sequences of the complementary strand of a polynucleotide sequence encoding an epitope of the invention, and polynucleotide sequences which hybridize to the complementary strand under stringent hybridization conditions or alternatively, under lower stringency hybridization conditions defined supra.

[0176] The term "epitopes," as used herein, refers to portions of a polypeptide having antigenic or immunogenic activity in an animal, preferably a mammal, and

most preferably in a human. In a preferred embodiment, the present invention encompasses a polypeptide comprising an epitope, as well as the polynucleotide encoding this polypeptide. An "immunogenic epitope," as used herein, is defined as a portion of a protein that elicits an antibody response in an animal, as determined by any method known in the art, for example, by the methods for generating antibodies described *infra*. (See, for example, Geysen et al., Proc. Natl. Acad. Sci. USA 81:3998- 4002 (1983)). The term "antigenic epitope," as used herein, is defined as a portion of a protein to which an antibody can immunospecifically bind its antigen as determined by any method well known in the art, for example, by the immunoassays described herein. Immunospecific binding excludes non-specific binding but does not necessarily exclude cross- reactivity with other antigens. Antigenic epitopes need not necessarily be immunogenic.

[0177] Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

In the present invention, antigenic epitopes preferably contain a sequence 101781 of at least 4, at least 5, at least 6, at least 7, more preferably at least 8, at least 9, at least 10, at least 11, at least 12, at least 13, at least 14, at least 15, at least 20, at least 25, at least 30, at least 40, at least 50, and, most preferably, between about 15 to about 30 amino acids. Preferred polypeptides comprising immunogenic or antigenic epitopes are at least 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, or 100 amino acid residues in length. Additional non-exclusive preferred antigenic epitopes include the antigenic epitopes disclosed herein, as well as portions thereof. Antigenic epitopes are useful, for example, to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. Preferred antigenic epitopes include the antigenic epitopes disclosed herein, as well as any combination of two, three, four, five or more of these antigenic epitopes. Antigenic epitopes can be used as the target molecules in immunoassays. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe et al., Science 219:660-666 (1983)).

[0179] Non-limiting examples of epitopes of polypeptides that can be used to generate antibodies of the invention include a polypeptide comprising, or alternatively consisting of, at least one, two, three, four, five, six or more of the portion(s) of SEQ

ID NO:Y specified in column 6 of Table 1A. These polypeptide fragments have been determined to bear antigenic epitopes of the proteins of the invention by the analysis of the Jameson-Wolf antigenic index which is included in the DNAStar suite of computer programs. By "comprise" it is intended that a polypeptide contains at least one, two, three, four, five, six or more of the portion(s) of SEQ ID NO:Y shown in column 6 of Table 1A, but it may contain additional flanking residues on either the amino or carboxyl termini of the recited portion. Such additional flanking sequences are preferably sequences naturally found adjacent to the portion; i.e., contiguous sequence shown in SEQ ID NO:Y. The flanking sequence may, however, be sequences from a heterologous polypeptide, such as from another protein described herein or from a heterologous polypeptide not described herein. In particular embodiments, epitope portions of a polypeptide of the invention comprise one, two, three, or more of the portions of SEQ ID NO:Y shown in column 6 of Table 1A. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0180] Similarly, immunogenic epitopes can be used, for example, to induce antibodies according to methods well known in the art. See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle et al., J. Gen. Virol. 66:2347-2354 (1985). Preferred immunogenic epitopes include the immunogenic epitopes disclosed herein, as well as any combination of two, three, four, five or more of these immunogenic epitopes. The polypeptides comprising one or more immunogenic epitopes may be presented for eliciting an antibody response together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse), or, if the polypeptide is of sufficient length (at least about 25 amino acids), the polypeptide may be presented without a carrier. However, immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting).

[0181] Epitope-bearing polypeptides of the present invention may be used to induce antibodies according to methods well known in the art including, but not limited to, in vivo immunization, in vitro immunization, and phage display methods. See, e.g., Sutcliffe et al., supra; Wilson et al., supra, and Bittle et al., J. Gen. Virol., 66:2347-2354 (1985). If in vivo immunization is used, animals may be immunized

with free peptide; however, anti-peptide antibody titer may be boosted by coupling the peptide to a macromolecular carrier, such as keyhole limpet hemacyanin (KLH) or tetanus toxoid. For instance, peptides containing cysteine residues may be coupled to a carrier using a linker such as maleimidobenzoyl- N-hydroxysuccinimide ester (MBS), while other peptides may be coupled to carriers using a more general linking agent such as glutaraldehyde. Animals such as rabbits, rats and mice are immunized with either free or carrier- coupled peptides, for instance, by intraperitoneal and/or intradermal injection of emulsions containing about 100 µg of peptide or carrier protein and Freund's adjuvant or any other adjuvant known for stimulating an immune response. Several booster injections may be needed, for instance, at intervals of about two weeks, to provide a useful titer of anti-peptide antibody which can be detected, for example, by ELISA assay using free peptide adsorbed to a solid surface. The titer of anti-peptide antibodies in serum from an immunized animal may be increased by selection of anti-peptide antibodies, for instance, by adsorption to the peptide on a solid support and elution of the selected antibodies according to methods well known in the art.

As one of skill in the art will appreciate, and as discussed above, the [0182]polypeptides of the present invention (e.g., those comprising an immunogenic or antigenic epitope) can be fused to heterologous polypeptide sequences. For example, polypeptides of the present invention (including fragments or variants thereof), may be fused with the constant domain of immunoglobulins (IgA, IgE, IgG, IgM), or portions thereof (CH1, CH2, CH3, or any combination thereof and portions thereof, resulting in chimeric polypeptides. By way of another non-limiting example, polypeptides and/or antibodies of the present invention (including fragments or variants thereof) may be fused with albumin (including but not limited to recombinant human serum albumin or fragments or variants thereof (see, e.g., U.S. Patent No. 5,876,969, issued March 2, 1999, EP Patent 0 413 622, and U.S. Patent No. 5,766,883, issued June 16, 1998, herein incorporated by reference in their entirety)). In a preferred embodiment, polypeptides and/or antibodies of the present invention (including fragments or variants thereof) are fused with the mature form of human serum albumin (i.e., amino acids 1 - 585 of human serum albumin as shown in Figures 1 and 2 of EP Patent 0 322 094) which is herein incorporated by reference in its entirety. In another preferred

embodiment, polypeptides and/or antibodies of the present invention (including fragments or variants thereof) are fused with polypeptide fragments comprising, or alternatively consisting of, amino acid residues 1-z of human serum albumin, where z is an integer from 369 to 419, as described in U.S. Patent 5,766,883 herein incorporated by reference in its entirety. Polypeptides and/or antibodies of the present invention (including fragments or variants thereof) may be fused to either the N- or C-terminal end of the heterologous protein (e.g., immunoglobulin Fc polypeptide or human serum albumin polypeptide). Polynucleotides encoding fusion proteins of the invention are also encompassed by the invention.

Such fusion proteins as those described above may facilitate purification [0183] and may increase half-life in vivo. This has been shown for chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. See, e.g., EP 394,827; Traunecker et al., Nature, 331:84-86 (1988). Enhanced delivery of an antigen across the epithelial barrier to the immune system has been demonstrated for antigens (e.g., insulin) conjugated to an FcRn binding partner such as IgG or Fc fragments (see, e.g., PCT Publications WO 96/22024 and WO 99/04813). IgG Fusion proteins that have a disulfide-linked dimeric structure due to the IgG portion desulfide bonds have also been found to be more efficient in binding and neutralizing other molecules than monomeric polypeptides or fragments thereof alone. See, e.g., Fountoulakis et al., J. Biochem., 270:3958-3964 (1995). Nucleic acids encoding the above epitopes can also be recombined with a gene of interest as an epitope tag (e.g., the hemagglutinin (iHAî) tag or flag tag) to aid in detection and purification of the expressed polypeptide. For example, a system described by Janknecht et al. allows for the ready purification of non-denatured fusion proteins expressed in human cell lines (Janknecht et al., 1991, Proc. Natl. Acad. Sci. USA 88:8972- 897). In this system, the gene of interest is subcloned into a vaccinia recombination plasmid such that the open reading frame of the gene is translationally fused to an amino-terminal tag consisting of six histidine residues. The tag serves as a matrix binding domain for the fusion protein. Extracts from cells infected with the recombinant vaccinia virus are loaded onto Ni2+ nitriloacetic acid-agarose column

and histidine-tagged proteins can be selectively eluted with imidazole-containing buffers.

Fusion Proteins

[0184] Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular locations based on trafficking signals, polypeptides of the present invention which are shown to be secreted can be used as targeting molecules once fused to other proteins.

[0185] Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

[0186] In certain preferred embodiments, proteins of the invention are fusion proteins comprising an amino acid sequence that is an N and/or C- terminal deletion of a polypeptide of the invention. In preferred embodiments, the invention is directed to a fusion protein comprising an amino acid sequence that is at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to a polypeptide sequence of the invention. Polynucleotides encoding these proteins are also encompassed by the invention.

[0187] Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell or subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to facilitate handling of polypeptides are familiar and routine techniques in the art.

[0188] As one of skill in the art will appreciate that, as discussed above, polypeptides of the present invention, and epitope-bearing fragments thereof, can be combined with heterologous polypeptide sequences. For example, the polypeptides of

the present invention may be fused with heterologous polypeptide sequences, for example, the polypeptides of the present invention may be fused with the constant domain of immunoglobulins (IgA, IgE, IgG, IgM) or portions thereof (CH1, CH2, CH3, and any combination thereof, including both entire domains and portions thereof), or albumin (including, but not limited to, native or recombinant human albumin or fragments or variants thereof (see, e.g., U.S. Patent No. 5,876,969, issued March 2, 1999, EP Patent 0 413 622, and U.S. Patent No. 5,766,883, issued June 16, 1998, herein incorporated by reference in their entirety)), resulting in chimeric polypeptides. For example, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties (EP-A 0232 262). Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. See, D. Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol. Chem. 270:9459-9471 (1995).

Moreover, the polypeptides of the present invention can be fused to marker sequences, such as a polypeptide which facilitates purification of the fused polypeptide. In preferred embodiments, the marker amino acid sequence is a hexahistidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein (Wilson et al., Cell 37:767 (1984).)

[0190] Additional fusion proteins of the invention may be generated through the techniques of gene-shuffling, motif-shuffling, exon-shuffling, and/or codon-shuffling

(collectively referred to as "DNA shuffling"), briefly described below, and further described herein. DNA shuffling may be employed to modulate the activities of polypeptides of the invention, such methods can be used to generate polypeptides with altered activity, as well as agonists and antagonists of the polypeptides. See, generally, U.S. Patent Nos. 5,605,793; 5,811,238; 5,830,721; 5,834,252; and 5,837,458, and Patten et al., Curr. Opinion Biotechnol. 8:724-33 (1997); Harayama, Trends Biotechnol. 16(2):76-82 (1998); Hansson et al., J. Mol. Biol. 287:265-76 (1999); and Lorenzo and Blasco, Biotechniques 24(2):308-13 (1998); each of these patents and publications are hereby incorporated by reference in its entirety). In a preferred embodiment, one or more components, motifs, sections, parts, domains, fragments, etc., of a polynucleotide encoding a polypeptide of the invention may be recombined with one or more components, motifs, sections, parts, domains, fragments, etc., of one or more heterologous molecules encoding a heterologous polypeptide.

[0191] Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the present invention.

Recombinant and Synthetic Production of Polypeptides of the Invention

[0192] The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by synthetic and recombinant techniques. The vector may be, for example, a phage, plasmid, viral, or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

[0193] The polynucleotides of the invention may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged *in vitro* using an appropriate packaging cell line and then transduced into host cells.

[0194] The polynucleotide insert should be operatively linked to an appropriate promoter, such as the phage lambda PL promoter, the *E. coli lac, trp, phoA* and *tac* promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to

name a few. Other suitable promoters will be known to the skilled artisan. The expression constructs will further contain sites for transcription initiation, termination, and, in the transcribed region, a ribosome binding site for translation. The coding portion of the transcripts expressed by the constructs will preferably include a translation initiating codon at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

[0195] As indicated, the expression vectors will preferably include at least one selectable marker. Such markers include dihydrofolate reductase, G418 or neomycin resistance, glutamine synthase, for eukaryotic cell culture and tetracycline, kanamycin or ampicillin resistance genes for culturing in *E. coli* and other bacteria. Representative examples of appropriate hosts include, but are not limited to, bacterial cells, such as *E. coli*, *Streptomyces* and *Salmonella typhimurium* cells; fungal cells, such as yeast cells (e.g., *Saccharomyces cerevisiae* or *Pichia pastoris* (ATCC Accession No. 201178)); insect cells such as *Drosophila* S2 and *Spodoptera* Sf9 cells; animal cells such as CHO, COS, 293, NSO and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

[0196] Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech, Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Preferred expression vectors for use in yeast systems include, but are not limited to pYES2, pYD1, pTEF1/Zeo, pYES2/GS, pPICZ, pGAPZ, pGAPZalph, pPIC9, pPIC3.5, pHIL-D2, pHIL-S1, pPIC3.5K, pPIC9K, and PAO815 (all available from Invitrogen, Carlsbad, CA). Other suitable vectors will be readily apparent to the skilled artisan.

[0197] Vectors which use glutamine synthase (GS) or DHFR as the selectable markers can be amplified in the presence of the drugs methionine sulphoximine or methotrexate, respectively. An advantage of glutamine synthase based vectors is the availabilty of cell lines (e.g., the murine myeloma cell line, NS0) which are glutamine

synthase negative. Glutamine synthase expression systems can also function in glutamine synthase expressing cells (e.g., Chinese Hamster Ovary (CHO) cells) by providing additional inhibitor to prevent the functioning of the endogenous gene. A glutamine synthase expression system and components thereof are detailed in PCT publications: WO87/04462; WO86/05807; WO89/01036; WO89/10404; and WO91/06657 which are hereby incorporated in their entireties by reference herein. Additionally, glutamine synthase expression vectors can be obtained from Lonza Biologics, Inc. (Portsmouth, NH). Expression and production of monoclonal antibodies using a GS expression system in murine myeloma cells is described in Bebbington et al., Bio/technology 10:169(1992) and in Biblia and Robinson Biotechnol. Prog. 11:1 (1995) which are herein incorporated by reference.

[0198] The present invention also relates to host cells containing the abovedescribed vector constructs described herein, and additionally encompasses host cells containing nucleotide sequences of the invention that are operably associated with one or more heterologous control regions (e.g., promoter and/or enhancer) using techniques known of in the art. The host cell can be a higher eukaryotic cell, such as a mammalian cell (e.g., a human derived cell), or a lower eukaryotic cell, such as a yeast cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. A host strain may be chosen which modulates the expression of the inserted gene sequences, or modifies and processes the gene product in the specific fashion desired. Expression from certain promoters can be elevated in the presence of certain inducers; thus expression of the genetically engineered polypeptide may be controlled. Furthermore, different host cells have characteristics and specific mechanisms for the translational and post-translational processing and modification (e.g., phosphorylation, cleavage) of proteins. Appropriate cell lines can be chosen to ensure the desired modifications and processing of the foreign protein expressed.

[0199] Introduction of the nucleic acids and nucleic acid constructs of the invention into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection, or other methods. Such methods are described in many standard laboratory manuals, such as Davis et al., Basic Methods In Molecular

Biology (1986). It is specifically contemplated that the polypeptides of the present invention may in fact be expressed by a host cell lacking a recombinant vector.

In addition to encompassing host cells containing the vector constructs [0200] discussed herein, the invention also encompasses primary, secondary, and immortalized host cells of vertebrate origin, particularly mammalian origin, that have been engineered to delete or replace endogenous genetic material (e.g., reproductive system antigen coding sequence), and/or to include genetic material (e.g., heterologous polynucleotide sequences) that is operably associated with reproductive system associated polynucleotides of the invention, and which activates, alters, and/or amplifies endogenous reproductive system associated polynucleotides. For example, techniques known in the art may be used to operably associate heterologous control regions (e.g., promoter and/or enhancer) and endogenous reproductive system associated polynucleotide sequences via homologous recombination (see, e.g., U.S. Patent Number 5,641,670, issued June 24, 1997; International Publication Number WO 96/29411; International Publication Number WO 94/12650; Koller et al., Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); and Zijlstra et al., Nature 342:435-438 (1989), the disclosures of each of which are incorporated by reference in their entireties).

Polypeptides of the present invention can also be recovered from: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes. Thus, it is well known in the art that the N-terminal methionine encoded by the translation initiation codon generally is removed with high efficiency from any protein after translation in all eukaryotic cells. While the N-terminal methionine on most proteins also is efficiently removed in most prokaryotes, for some proteins, this prokaryotic removal process is inefficient,

depending on the nature of the amino acid to which the N-terminal methionine is covalently linked.

[0202] In one embodiment, the yeast Pichia pastoris is used to express polypeptides of the invention in a eukaryotic system. Pichia pastoris is a methylotrophic yeast which can metabolize methanol as its sole carbon source. A main step in the methanol metabolization pathway is the oxidation of methanol to formaldehyde using O2. This reaction is catalyzed by the enzyme alcohol oxidase. In order to metabolize methanol as its sole carbon source, Pichia pastoris must generate high levels of alcohol oxidase due, in part, to the relatively low affinity of alcohol oxidase for O2. Consequently, in a growth medium depending on methanol as a main carbon source, the promoter region of one of the two alcohol oxidase genes (AOXI) is highly active. In the presence of methanol, alcohol oxidase produced from the AOX1 gene comprises up to approximately 30% of the total soluble protein in Pichia pastoris. See, Ellis, S.B., et al., Mol. Cell. Biol. 5:1111-21 (1985); Koutz, P.J, et al., Yeast 5:167-77 (1989); Tschopp, J.F., et al., Nucl. Acids Res. 15:3859-76 (1987). Thus, a heterologous coding sequence, such as, for example, a polynucleotide of the present invention, under the transcriptional regulation of all or part of the AOXI regulatory sequence is expressed at exceptionally high levels in Pichia yeast grown in the presence of methanol.

[0203] In one example, the plasmid vector pPIC9K is used to express DNA encoding a polypeptide of the invention, as set forth herein, in a *Pichea* yeast system essentially as described in "*Pichia* Protocols: Methods in Molecular Biology," D.R. Higgins and J. Cregg, eds. The Humana Press, Totowa, NJ, 1998. This expression vector allows expression and secretion of a polypeptide of the invention by virtue of the strong *AOX1* promoter linked to the *Pichia pastoris* alkaline phosphatase (PHO) secretory signal peptide (i.e., leader) located upstream of a multiple cloning site.

[0204] Many other yeast vectors could be used in place of pPIC9K, such as, pYES2, pYD1, pTEF1/Zeo, pYES2/GS, pPICZ, pGAPZ, pGAPZalpha, pPIC9, pPIC3.5, pHIL-D2, pHIL-S1, pPIC3.5K, and PAO815, as one skilled in the art would readily appreciate, as long as the proposed expression construct provides appropriately

located signals for transcription, translation, secretion (if desired), and the like, including an in-frame AUG as required.

[0205] In another embodiment, high-level expression of a heterologous coding sequence, such as, for example, a polynucleotide of the present invention, may be achieved by cloning the heterologous polynucleotide of the invention into an expression vector such as, for example, pGAPZ or pGAPZalpha, and growing the yeast culture in the absence of methanol.

In addition to encompassing host cells containing the vector constructs [0206] discussed herein, the invention also encompasses primary, secondary, and immortalized host cells of vertebrate origin, particularly mammalian origin, that have been engineered to delete or replace endogenous genetic material (e.g., coding sequence), and/or to include genetic material (e.g., heterologous polynucleotide sequences) that is operably associated with polynucleotides of the invention, and which activates, alters, and/or amplifies endogenous polynucleotides. For example, techniques known in the art may be used to operably associate heterologous control regions (e.g., promoter and/or enhancer) and endogenous polynucleotide sequences via homologous recombination (see, e.g., U.S. Patent No. 5,641,670, issued June 24, 1997; International Publication No. WO 96/29411, published September 26, 1996; International Publication No. WO 94/12650, published August 4, 1994; Koller et al., Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); and Zijlstra et al., Nature 342:435-438 (1989), the disclosures of each of which are incorporated by reference in their entireties).

In addition, polypeptides of the invention can be chemically synthesized using techniques known in the art (e.g., see Creighton, 1983, Proteins: Structures and Molecular Principles, W.H. Freeman & Co., N.Y., and Hunkapiller et al., Nature, 310:105-111 (1984)). For example, a polypeptide corresponding to a fragment of a polypeptide can be synthesized by use of a peptide synthesizer. Furthermore, if desired, nonclassical amino acids or chemical amino acid analogs can be introduced as a substitution or addition into the polypeptide sequence. Non-classical amino acids include, but are not limited to, to the D-isomers of the common amino acids, 2,4-diaminobutyric acid, a-amino isobutyric acid, 4-aminobutyric acid, Abu, 2-amino butyric acid, g-Abu, e-Ahx, 6-amino hexanoic acid, Aib, 2-amino isobutyric acid,

3-amino propionic acid, omithine, norleucine, norvaline, hydroxyproline, sarcosine, citrulline, homocitrulline, cysteic acid, t-butylglycine, t-butylalanine, phenylglycine, cyclohexylalanine, b-alanine, fluoro-amino acids, designer amino acids such as b-methyl amino acids, Ca-methyl amino acids, Na-methyl amino acids, and amino acid analogs in general. Furthermore, the amino acid can be D (dextrorotary) or L (levorotary).

[0208] The invention encompasses polypeptides of the present invention which are differentially modified during or after translation, e.g., by glycosylation, acetylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to an antibody molecule or other cellular ligand, etc. Any of numerous chemical modifications may be carried out by known techniques, including but not limited, to specific chemical cleavage by cyanogen bromide, trypsin, chymotrypsin, papain, V8 protease, NaBH₄; acetylation, formylation, oxidation, reduction; metabolic synthesis in the presence of tunicamycin; etc.

[0209] Additional post-translational modifications encompassed by the invention include, for example, e.g., N-linked or O-linked carbohydrate chains, processing of N-terminal or C-terminal ends), attachment of chemical moieties to the amino acid backbone, chemical modifications of N-linked or O-linked carbohydrate chains, and addition or deletion of an N-terminal methionine residue as a result of procaryotic host cell expression. The polypeptides may also be modified with a detectable label, such as an enzymatic, fluorescent, isotopic or affinity label to allow for detection and isolation of the protein.

[0210] Examples of suitable enzymes include horseradish peroxidase, alkaline phosphatase, beta-galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin; and examples of suitable radioactive material include iodine (121 I, 123 I, 125 I, 131 I), carbon (14 C), sulfur (35 S), tritium (3H), indium (111 In, 112 In, 113 m In, 115 m In), technetium (99 Tc, 99 m Tc), thallium (201 Ti), gallium (68 Ga, 67 Ga), palladium (103 Pd), molybdenum (99 Mo), xenon

(133Xe), fluorine (18F), 153Sm; 177Lu, 159Gd, 149Pm, 140La, 175Yb, 166Ho, 90Y, 47Sc, 186Re, 188Re, 142Pr, 105Rh, and 97Ru.

In specific embodiments, a polypeptide of the present invention or fragment or variant thereof is attached to macrocyclic chelators that associate with radiometal ions, including but not limited to, ¹⁷⁷Lu, ⁹⁰Y, ¹⁶⁶Ho, and ¹⁵³Sm, to polypeptides. In a preferred embodiment, the radiometal ion associated with the macrocyclic chelators is ¹¹¹In. In another preferred embodiment, the radiometal ion associated with the macrocyclic chelator is ⁹⁰Y. In specific embodiments, the macrocyclic chelator is 1,4,7,10-tetraazacyclododecane-N,N',N",N"'-tetraacetic acid (DOTA). In other specific embodiments, DOTA is attached to an antibody of the invention or fragment thereof via a linker molecule. Examples of linker molecules useful for conjugating DOTA to a polypeptide are commonly known in the art - see, for example, DeNardo et al., Clin Cancer Res. 4(10):2483-90 (1998); Peterson et al., Bioconjug. Chem. 10(4):553-7 (1999); and Zimmerman et al, Nucl. Med. Biol. 26(8):943-50 (1999); which are hereby incorporated by reference in their entirety.

As mentioned, the reproductive system associated proteins of the invention [0212] may be modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given reproductive system associated polypeptide. Reproductive system associated polypeptides may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic reproductive system associated polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphotidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization,

selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth. Enzymol. 182:626-646 (1990); Rattan et al., Ann. N.Y. Acad. Sci. 663:48-62 (1992)).

[0213] Also provided by the invention are chemically modified derivatives of the polypeptides of the invention which may provide additional advantages such as increased solubility, stability and circulating time of the polypeptide, or decreased immunogenicity (see U.S. Patent No. 4,179,337). The chemical moieties for derivitization may be selected from water soluble polymers such as polyethylene glycol, ethylene glycol/propylene glycol copolymers, carboxymethylcellulose, dextran, polyvinyl alcohol and the like. The polypeptides may be modified at random positions within the molecule, or at predetermined positions within the molecule and may include one, two, three or more attached chemical mojeties.

The polymer may be of any molecular weight, and may be branched or [0214] unbranched. For polyethylene glycol, the preferred molecular weight is between about 1 kDa and about 100 kDa (the term "about" indicating that in preparations of polyethylene glycol, some molecules will weigh more, some less, than the stated molecular weight) for ease in handling and manufacturing. Other sizes may be used, depending on the desired therapeutic profile (e.g., the duration of sustained release desired, the effects, if any on biological activity, the ease in handling, the degree or lack of antigenicity and other known effects of the polyethylene glycol to a therapeutic protein or analog). For example, the polyethylene glycol may have an average molecular weight of about 200, 500, 1000, 1500, 2000, 2500, 3000, 3500, 4000, 4500, 5000, 5500, 6000, 6500, 7000, 7500, 8000, 8500, 9000, 9500, 10,000, 10,500, 11,000, 11,500, 12,000, 12,500, 13,000, 13,500, 14,000, 14,500, 15,000, 15,500, 16,000, 16,500, 17,000, 17,500, 18,000, 18,500, 19,000, 19,500, 20,000, 25,000, 30,000, 35,000, 40,000, 50,000, 55,000, 60,000, 65,000, 70,000, 75,000, 80,000, 85,000, 90,000, 95,000, or 100,000 kDa.

[0215] As noted above, the polyethylene glycol may have a branched structure.

Branched polyethylene glycols are described, for example, in U.S. Patent No. 5,643,575; Morpurgo et al., Appl. Biochem. Biotechnol. 56:59-72 (1996); Vorobjev et al., Nucleosides Nucleotides 18:2745-2750 (1999); and Caliceti et al., Bioconjug. Chem. 10:638-646 (1999), the disclosures of each of which are incorporated herein by reference.

[0216] The polyethylene glycol molecules (or other chemical moieties) should be attached to the protein with consideration of effects on functional or antigenic domains of the protein. There are a number of attachment methods available to those skilled in the art, such as, for example, the method disclosed in EP 0 401 384 (coupling PEG to G-CSF), herein incorporated by reference; see also Malik et al., Exp. Hematol. 20:1028-1035 (1992), reporting pegylation of GM-CSF using tresyl chloride. For example, polyethylene glycol may be covalently bound through amino acid residues via a reactive group, such as a free amino or carboxyl group. Reactive groups are those to which an activated polyethylene glycol molecule may be bound. The amino acid residues having a free amino group may include lysine residues and the N-terminal amino acid residues; those having a free carboxyl group may include aspartic acid residues glutamic acid residues and the C-terminal amino acid residue. Sulfhydryl groups may also be used as a reactive group for attaching the polyethylene glycol molecules. Preferred for therapeutic purposes is attachment at an amino group, such as attachment at the N-terminus or lysine group.

As suggested above, polyethylene glycol may be attached to proteins via linkage to any of a number of amino acid residues. For example, polyethylene glycol can be linked to proteins via covalent bonds to lysine, histidine, aspartic acid, glutamic acid, or cysteine residues. One or more reaction chemistries may be employed to attach polyethylene glycol to specific amino acid residues (e.g., lysine, histidine, aspartic acid, glutamic acid, or cysteine) of the protein or to more than one type of amino acid residue (e.g., lysine, histidine, aspartic acid, glutamic acid, cysteine and combinations thereof) of the protein.

[0218] One may specifically desire proteins chemically modified at the N-terminus. Using polyethylene glycol as an illustration of the present composition, one may select from a variety of polyethylene glycol molecules (by molecular weight, branching, etc.), the proportion of polyethylene glycol molecules to protein

(polypeptide) molecules in the reaction mix, the type of pegylation reaction to be performed, and the method of obtaining the selected N-terminally pegylated protein. The method of obtaining the N-terminally pegylated preparation (i.e., separating this moiety from other monopegylated moieties if necessary) may be by purification of the N-terminally pegylated material from a population of pegylated protein molecules. Selective proteins chemically modified at the N-terminus modification may be accomplished by reductive alkylation which exploits differential reactivity of different types of primary amino groups (lysine versus the N-terminal) available for derivatization in a particular protein. Under the appropriate reaction conditions, substantially selective derivatization of the protein at the N-terminus with a carbonyl group containing polymer is achieved.

[0219] As indicated above, pegylation of the proteins of the invention may be accomplished by any number of means. For example, polyethylene glycol may be attached to the protein either directly or by an intervening linker. Linkerless systems for attaching polyethylene glycol to proteins are described in Delgado et al., Crit. Rev. Thera. Drug Carrier Sys. 9:249-304 (1992); Francis et al., Intern. J. of Hematol. 68:1-18 (1998); U.S. Patent No. 4,002,531; U.S. Patent No. 5,349,052; WO 95/06058; and WO 98/32466, the disclosures of each of which are incorporated herein by reference.

[0220] One system for attaching polyethylene glycol directly to amino acid residues of proteins without an intervening linker employs tresylated MPEG, which is produced by the modification of monmethoxy polyethylene glycol (MPEG) using tresylchloride (CISO₂CH₂CF₃). Upon reaction of protein with tresylated MPEG, polyethylene glycol is directly attached to amine groups of the protein. Thus, the invention includes protein-polyethylene glycol conjugates produced by reacting proteins of the invention with a polyethylene glycol molecule having a 2,2,2-trifluoreothane sulphonyl group.

[0221] Polyethylene glycol can also be attached to proteins using a number of different intervening linkers. For example, U.S. Patent No. 5,612,460, the entire disclosure of which is incorporated herein by reference, discloses urethane linkers for connecting polyethylene glycol to proteins. Protein-polyethylene glycol conjugates wherein the polyethylene glycol is attached to the protein by a linker can also be produced by reaction of proteins with compounds such as MPEG-

succinimidylsuccinate, MPEG activated with 1,1'-carbonyldiimidazole, MPEG-2,4,5-trichloropenylcarbonate, MPEG-p-nitrophenolcarbonate, and various MPEG-succinate derivatives. A number of additional polyethylene glycol derivatives and reaction chemistries for attaching polyethylene glycol to proteins are described in International Publication No. WO 98/32466, the entire disclosure of which is incorporated herein by reference. Pegylated protein products produced using the reaction chemistries set out herein are included within the scope of the invention.

The number of polyethylene glycol moieties attached to each protein of the invention (i.e., the degree of substitution) may also vary. For example, the pegylated proteins of the invention may be linked, on average, to 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 12, 15, 17, 20, or more polyethylene glycol molecules. Similarly, the average degree of substitution within ranges such as 1-3, 2-4, 3-5, 4-6, 5-7, 6-8, 7-9, 8-10, 9-11, 10-12, 11-13, 12-14, 13-15, 14-16, 15-17, 16-18, 17-19, or 18-20 polyethylene glycol moieties per protein molecule. Methods for determining the degree of substitution are discussed, for example, in Delgado et al., Crit. Rev. Thera. Drug Carrier Sys. 9:249-304 (1992).

The reproductive system associated polypeptides of the invention can be recovered and purified from chemical synthesis and recombinant cell cultures by standard methods which include, but are not limited to, ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification. Well known techniques for refolding protein may be employed to regenerate active conformation when the polypeptide is denatured during isolation and/or purification.

[0224] Reproductive system associated polynucleotides and polypeptides may be used in accordance with the present invention for a variety of applications, particularly those that make use of the chemical and biological properties of reproductive system associated antigens. Among these are applications in the detection, prevention, diagnosis and/or treatment of diseases associated with the reproductive system, such as e.g., cancers of the reproductive system, tumors, injuries and trauma, infections,

congenital defects, and diseases or disorders which result in infertility, complications with pregnancy, labor, or parturition, postpartum difficulties, and as listed below in the section entitled "Reproductive System Disorders". Additional applications relate to diagnosis and to treatment of disorders of cells, tissues and organisms. These aspects of the invention are discussed further below.

[0225] In a preferred embodiment, polynucleotides expressed in a particular tissue type (see, e.g., Table 1A, column 7) are used to detect, diagnose, treat, prevent and/or prognose disorders associated with the tissue type.

[0226] The polypeptides of the invention may be in monomers or multimers (i.e., dimers, trimers, tetramers and higher multimers). Accordingly, the present invention relates to monomers and multimers of the polypeptides of the invention, their preparation, and compositions (preferably, Therapeutics) containing them. In specific embodiments, the polypeptides of the invention are monomers, dimers, trimers or tetramers. In additional embodiments, the multimers of the invention are at least dimers, at least trimers, or at least tetramers.

Multimers encompassed by the invention may be homomers or heteromers. [0227] As used herein, the term homomer refers to a multimer containing only polypeptides corresponding to a protein of the invention (e.g., the amino acid sequence of SEQ ID NO:Y, an amino acid sequence encoded by SEQ ID NO:X or the complement of SEQ ID NO:X, the amino acid sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or an amino acid sequence encoded by cDNA contained in Clone ID NO:Z (including fragments, variants, splice variants, and fusion proteins, corresponding to these as described herein)). These homomers may contain polypeptides having identical or different amino acid sequences. In a specific embodiment, a homomer of the invention is a multimer containing only polypeptides having an identical amino acid sequence. In another specific embodiment, a homomer of the invention is a multimer containing polypeptides having different amino acid sequences. In specific embodiments, the multimer of the invention is a homodimer (e.g., containing two polypeptides having identical or different amino acid sequences) or a homotrimer (e.g., containing three polypeptides having identical and/or different amino acid sequences). In additional embodiments, the homomeric multimer of the invention is at least a homodimer, at least a homotrimer, or at least a homotetramer.

[0228] As used herein, the term heteromer refers to a multimer containing two or more heterologous polypeptides (i.e., polypeptides of different proteins) in addition to the polypeptides of the invention. In a specific embodiment, the multimer of the invention is a heterodimer, a heterotrimer, or a heterotetramer. In additional embodiments, the heteromeric multimer of the invention is at least a heterodimer, at least a heterotrimer, or at least a heterotetramer.

Multimers of the invention may be the result of hydrophobic, hydrophilic, [0229] ionic and/or covalent associations and/or may be indirectly linked by, for example, liposome formation. Thus, in one embodiment, multimers of the invention, such as, for example, homodimers or homotrimers, are formed when polypeptides of the invention contact one another in solution. In another embodiment, heteromultimers of the invention, such as, for example, heterotrimers or heterotetramers, are formed when polypeptides of the invention contact antibodies to the polypeptides of the invention (including antibodies to the heterologous polypeptide sequence in a fusion protein of the invention) in solution. In other embodiments, multimers of the invention are formed by covalent associations with and/or between the polypeptides of the invention. Such covalent associations may involve one or more amino acid residues contained in the polypeptide sequence (e.g., that recited in SEQ ID NO:Y, encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or encoded by the cDNA contained in Clone ID NO:Z). In one instance, the covalent associations are cross-linking between cysteine residues located within the polypeptide sequences which interact in the native (i.e., naturally occurring) polypeptide. In another instance, the covalent associations are the consequence of chemical or recombinant manipulation. Alternatively, such covalent associations may involve one or more amino acid residues contained in the heterologous polypeptide sequence in a fusion protein. In one example, covalent associations are between the heterologous sequence contained in a fusion protein of the invention (see, e.g., U.S. Patent Number 5,478,925). In a specific example, the covalent associations are between the heterologous sequence contained in a Fc fusion protein of the invention (as described herein). In another specific example, covalent associations of fusion proteins of the invention are between heterologous polypeptide sequence from another protein that is capable of forming covalently associated multimers, such as for example,

osteoprotegerin (see, e.g., International Publication NO: WO 98/49305, the contents of which are herein incorporated by reference in its entirety). In another embodiment, two or more polypeptides of the invention are joined through peptide linkers. Examples include those peptide linkers described in U.S. Pat. No. 5,073,627 (hereby incorporated by reference). Proteins comprising multiple polypeptides of the invention separated by peptide linkers may be produced using conventional recombinant DNA technology.

[0230] Another method for preparing multimer polypeptides of the invention involves use of polypeptides of the invention fused to a leucine zipper or isoleucine zipper polypeptide sequence. Leucine zipper and isoleucine zipper domains are polypeptides that promote multimerization of the proteins in which they are found. Leucine zippers were originally identified in several DNA-binding proteins (Landschulz et al., Science 240:1759, (1988)), and have since been found in a variety of different proteins. Among the known leucine zippers are naturally occurring peptides and derivatives thereof that dimerize or trimerize. Examples of leucine zipper domains suitable for producing soluble multimeric proteins of the invention are those described in PCT application WO 94/10308, hereby incorporated by reference. Recombinant fusion proteins comprising a polypeptide of the invention fused to a polypeptide sequence that dimerizes or trimerizes in solution are expressed in suitable host cells, and the resulting soluble multimeric fusion protein is recovered from the culture supernatant using techniques known in the art.

[0231] Trimeric polypeptides of the invention may offer the advantage of enhanced biological activity. Preferred leucine zipper moieties and isoleucine moieties are those that preferentially form trimers. One example is a leucine zipper derived from lung surfactant protein D (SPD), as described in Hoppe et al. (FEBS Letters 344:191, (1994)) and in U.S. patent application Ser. No. 08/446,922, hereby incorporated by reference. Other peptides derived from naturally occurring trimeric proteins may be employed in preparing trimeric polypeptides of the invention.

[0232] In another example, proteins of the invention are associated by interactions between Flag® polypeptide sequence contained in fusion proteins of the invention containing Flag® polypeptide sequence. In a further embodiment, proteins of the

invention are associated by interactions between heterologous polypeptide sequence contained in Flag® fusion proteins of the invention and anti-Flag® antibody.

The multimers of the invention may be generated using chemical [0233] techniques known in the art. For example, polypeptides desired to be contained in the multimers of the invention may be chemically cross-linked using linker molecules and linker molecule length optimization techniques known in the art (see, e.g., U.S. Patent Number 5,478,925, which is herein incorporated by reference in its entirety). Additionally, multimers of the invention may be generated using techniques known in the art to form one or more inter-molecule cross-links between the cysteine residues located within the sequence of the polypeptides desired to be contained in the multimer (see, e.g., U.S. Patent Number 5,478,925, which is herein incorporated by reference in its entirety). Further, polypeptides of the invention may be routinely modified by the addition of cysteine or biotin to the C-terminus or N-terminus of the polypeptide and techniques known in the art may be applied to generate multimers containing one or more of these modified polypeptides (see, e.g., U.S. Patent Number 5,478,925, which is herein incorporated by reference in its entirety). Additionally, techniques known in the art may be applied to generate liposomes containing the polypeptide components desired to be contained in the multimer of the invention (see, e.g., U.S. Patent Number 5,478,925, which is herein incorporated by reference in its entirety).

[0234] Alternatively, multimers of the invention may be generated using genetic engineering techniques known in the art. In one embodiment, polypeptides contained in multimers of the invention are produced recombinantly using fusion protein technology described herein or otherwise known in the art (see, e.g., U.S. Patent Number 5,478,925, which is herein incorporated by reference in its entirety). In a specific embodiment, polynucleotides coding for a homodimer of the invention are generated by ligating a polynucleotide sequence encoding a polypeptide of the invention to a sequence encoding a linker polypeptide and then further to a synthetic polynucleotide encoding the translated product of the polypeptide in the reverse orientation from the original C-terminus to the N-terminus (lacking the leader sequence) (see, e.g., U.S Patent Number 5,478,925, which is herein incorporated by reference in its entirety). In another embodiment, recombinant techniques described

herein or otherwise known in the art are applied to generate recombinant polypeptides of the invention which contain a transmembrane domain (or hydrophobic or signal peptide) and which can be incorporated by membrane reconstitution techniques into liposomes (see, e.g., U.S. Patent Number 5,478,925, which is herein incorporated by reference in its entirety).

Antibodies

[0235] Further polypeptides of the invention relate to antibodies and T-cell antigen receptors (TCR) which immunospecifically bind a polypeptide, polypeptide fragment, or variant of the invention (e.g., a polypeptide or fragment or variant of the amino acid sequence of SEQ ID NO:Y or a polypeptide encoded by the cDNA contained in Clone ID NO:Z, and/or an epitope, of the present invention) as determined by immunoassays well known in the art for assaying specific antibody-antigen binding. Antibodies of the invention include, but are not limited to, polyclonal, monoclonal, multispecific, human, humanized or chimeric antibodies, single chain antibodies, Fab fragments, F(ab') fragments, fragments produced by a Fab expression library, anti-idiotypic (anti-Id) antibodies (including, e.g., anti-Id antibodies to antibodies of the invention), intracellularly-made antibodies (i.e., intrabodies), and epitope-binding fragments of any of the above. The term "antibody," as used herein, refers to immunoglobulin molecules and immunologically active portions of immunoglobulin molecules, i.e., molecules that contain an antigen binding site that immunospecifically binds an antigen. The immunoglobulin molecules of the invention can be of any type (e.g., IgG, IgE, IgM, IgD, IgA and IgY), class (e.g., IgG1, IgG2, IgG3, IgG4, IgA1 and IgA2) or subclass of immunoglobulin molecule. In preferred embodiments, the immunoglobulin molecules of the invention are IgG1. In other preferred embodiments, the immunoglobulin molecules of the invention are IgG4.

[0236] Most preferably the antibodies are human antigen-binding antibody fragments of the present invention and include, but are not limited to, Fab, Fab' and F(ab')2, Fd, single-chain Fvs (scFv), single-chain antibodies, disulfide-linked Fvs (sdFv) and fragments comprising either a VL or VH domain. Antigen-binding antibody fragments, including single-chain antibodies, may comprise the variable region(s) alone or in combination with the entirety or a portion of the following: hinge

region, CH1, CH2, and CH3 domains. Also included in the invention are antigen-binding fragments also comprising any combination of variable region(s) with a hinge region, CH1, CH2, and CH3 domains. The antibodies of the invention may be from any animal origin including birds and mammals. Preferably, the antibodies are human, murine (e.g., mouse and rat), donkey, ship rabbit, goat, guinea pig, camel, horse, or chicken. As used herein, "human" antibodies include antibodies having the amino acid sequence of a human immunoglobulin and include antibodies isolated from human immunoglobulin libraries or from animals transgenic for one or more human immunoglobulin and that do not express endogenous immunoglobulins, as described infra and, for example in, U.S. Patent No. 5,939,598 by Kucherlapati et al.

The antibodies of the present invention may be monospecific, bispecific, trispecific or of greater multispecificity. Multispecific antibodies may be specific for different epitopes of a polypeptide of the present invention or may be specific for both a polypeptide of the present invention as well as for a heterologous epitope, such as a heterologous polypeptide or solid support material. See, e.g., PCT publications WO 93/17715; WO 92/08802; WO 91/00360; WO 92/05793; Tutt, et al., J. Immunol. 147:60-69 (1991); U.S. Patent Nos. 4,474,893; 4,714,681; 4,925,648; 5,573,920; 5,601,819; Kostelny et al., J. Immunol. 148:1547-1553 (1992).

of the epitope(s) or portion(s) of a polypeptide of the present invention which they recognize or specifically bind. The epitope(s) or polypeptide portion(s) may be specified as described herein, e.g., by N-terminal and C-terminal positions, or by size in contiguous amino acid residues, or listed in the Tables and Figures. Preferred epitopes of the invention include those shown in column 6 of Table 1A, as well as polynucleotides that encode these epitopes. Antibodies which specifically bind any epitope or polypeptide of the present invention may also be excluded. Therefore, the present invention includes antibodies that specifically bind polypeptides of the present invention, and allows for the exclusion of the same.

[0239] Antibodies of the present invention may also be described or specified in terms of their cross-reactivity. Antibodies that do not bind any other analog, ortholog, or homolog of a polypeptide of the present invention are included. Antibodies that bind polypeptides with at least 95%, at least 90%, at least 85%, at least 80%, at least

75%, at least 70%, at least 65%, at least 60%, at least 55%, and at least 50% identity (as calculated using methods known in the art and described herein) to a polypeptide of the present invention are also included in the present invention. In specific embodiments, antibodies of the present invention cross-react with murine, rat and/or rabbit homologs of human proteins and the corresponding epitopes thereof. Antibodies that do not bind polypeptides with less than 95%, less than 90%, less than 85%, less than 80%, less than 75%, less than 70%, less than 65%, less than 60%, less than 55%, and less than 50% identity (as calculated using methods known in the art and described herein) to a polypeptide of the present invention are also included in the present invention. In a specific embodiment, the above-described cross-reactivity is with respect to any single specific antigenic or immunogenic polypeptide, or combination(s) of 2, 3, 4, 5, or more of the specific antigenic and/or immunogenic polypeptides disclosed herein. Further included in the present invention are antibodies which bind polypeptides encoded by polynucleotides which hybridize to a polynucleotide of the present invention under stringent hybridization conditions (as described herein). Antibodies of the present invention may also be described or specified in terms of their binding affinity to a polypeptide of the invention. Preferred binding affinities include those with a dissociation constant or Kd less than 5 X 10⁻² M, 10^{-2} M, 5 X 10^{-3} M, 10^{-3} M, 5 X 10^{-4} M, 10^{-4} M, 5 X 10^{-5} M, 10^{-5} M, 5 X 10^{-6} M, 10^{-6} M, 5 X 10^{-7} M, 10^{7} M, 5 X 10^{-8} M, 10^{-8} M, 5 X 10^{-9} M, 10^{-9} M, 5 X 10^{-10} M, 10^{-10} M, 5 X 10^{-11} M, 10^{-11} M, 5 X 10^{-12} M, 10^{-12} M, 5 X 10^{-13} M, 10^{-13} M, 5 X 10^{-14} M, 10^{-14} 14 M, 5 X 10^{-15} M, or 10^{-15} M.

[0240] The invention also provides antibodies that competitively inhibit binding of an antibody to an epitope of the invention as determined by any method known in the art for determining competitive binding, for example, the immunoassays described herei-n. In preferred embodiments, the antibody competitively inhibits binding to the epitope by at least 95%, at least 90%, at least 85%, at least 80%, at least 75%, at least 70%, at least 60%, or at least 50%.

[0241] Antibodies of the present invention may act as agonists or antagonists of the polypeptides of the present invention. For example, the present invention includes antibodies which disrupt the receptor/ligand interactions with the polypeptides of the invention either partially or fully. Preferably, antibodies of the present invention bind

an antigenic epitope disclosed herein, or a portion thereof. The invention features both receptor-specific antibodies and ligand-specific antibodies. The invention also features receptor-specific antibodies which do not prevent ligand binding but prevent receptor activation. Receptor activation (i.e., signaling) may be determined by techniques described herein or otherwise known in the art. For example, receptor activation can be determined by detecting the phosphorylation (e.g., tyrosine or serine/threonine) of the receptor or its substrate by immunoprecipitation followed by western blot analysis (for example, as described *supra*). In specific embodiments, antibodies are provided that inhibit ligand activity or receptor activity by at least 95%, at least 90%, at least 85%, at least 80%, at least 75%, at least 70%, at least 60%, or at least 50% of the activity in absence of the antibody.

[0242] The invention also features receptor-specific antibodies which both prevent ligand binding and receptor activation as well as antibodies that recognize the receptor-ligand complex, and, preferably, do not specifically recognize the unbound receptor or the unbound ligand. Likewise, included in the invention are neutralizing antibodies which bind the ligand and prevent binding of the ligand to the receptor, as well as antibodies which bind the ligand, thereby preventing receptor activation, but do not prevent the ligand from binding the receptor. Further included in the invention are antibodies which activate the receptor. These antibodies may act as receptor agonists, i.e., potentiate or activate either all or a subset of the biological activities of the ligand-mediated receptor activation, for example, by inducing dimerization of the receptor. The antibodies may be specified as agonists, antagonists or inverse agonists for biological activities comprising the specific biological activities of the peptides of the invention disclosed herein. The above antibody agonists can be made using methods known in the art. See, e.g., PCT publication WO 96/40281; U.S. Patent No. 5,811,097; Deng et al., Blood 92(6):1981-1988 (1998); Chen et al., Cancer Res. 58(16):3668-3678 (1998); Harrop et al., J. Immunol. 161(4):1786-1794 (1998); Zhu et al., Cancer Res. 58(15):3209-3214 (1998); Yoon et al., J. Immunol. 160(7):3170-3179 (1998); Prat et al., J. Cell. Sci. 111(Pt2):237-247 (1998); Pitard et al., J. Immunol. Methods 205(2):177-190 (1997); Liautard et al., Cytokine 9(4):233-241 (1997); Carlson et al., J. Biol. Chem. 272(17):11295-11301 (1997); Taryman et al., Neuron 14(4):755-762 (1995); Muller et al., Structure 6(9):1153-1167 (1998); Bartunek et al.,

Cytokine 8(1):14-20 (1996) (which are all incorporated by reference herein in their entireties).

[0243] Antibodies of the present invention may be used, for example, to purify, detect, and target the polypeptides of the present invention, including both in vitro and in vivo diagnostic and therapeutic methods. For example, the antibodies have utility in immunoassays for qualitatively and quantitatively measuring levels of the polypeptides of the present invention in biological samples. See, e.g., Harlow et al., Antibodies: A Laboratory Manual, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988); incorporated by reference herein in its entirety.

As discussed in more detail below, the antibodies of the present invention may be used either alone or in combination with other compositions. The antibodies may further be recombinantly fused to a heterologous polypeptide at the N- or C-terminus or chemically conjugated (including covalent and non-covalent conjugations) to polypeptides or other compositions. For example, antibodies of the present invention may be recombinantly fused or conjugated to molecules useful as labels in detection assays and effector molecules such as heterologous polypeptides, drugs, radionuclides, or toxins. See, e.g., PCT publications WO 92/08495; WO 91/14438; WO 89/12624; U.S. Patent No. 5,314,995; and EP 396,387; the disclosures of which are incorporated herein by reference in their entireties.

[0245] The antibodies of the invention include derivatives that are modified, i.e., by the covalent attachment of any type of molecule to the antibody such that covalent attachment does not prevent the antibody from generating an anti-idiotypic response. For example, but not by way of limitation, the antibody derivatives include antibodies that have been modified, e.g., by glycosylation, acetylation, pegylation, phosphylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to a cellular ligand or other protein, etc. Any of numerous chemical modifications may be carried out by known techniques, including, but not limited to specific chemical cleavage, acetylation, formylation, metabolic synthesis of tunicamycin, etc. Additionally, the derivative may contain one or more non-classical amino acids.

[0246] The antibodies of the present invention may be generated by any suitable method known in the art. Polyclonal antibodies to an antigen-of- interest can be

produced by various procedures well known in the art. For example, a polypeptide of the invention can be administered to various host animals including, but not limited to, rabbits, mice, rats, etc. to induce the production of sera containing polyclonal antibodies specific for the antigen. Various adjuvants may be used to increase the immunological response, depending on the host species, and include but are not limited to, Freund's (complete and incomplete), mineral gels such as aluminum hydroxide, surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanins, dinitrophenol, and potentially useful human adjuvants such as BCG (bacille Calmette-Guerin) and corynebacterium parvum. Such adjuvants are also well known in the art.

Monoclonal antibodies can be prepared using a wide variety of techniques known in the art including the use of hybridoma, recombinant, and phage display technologies, or a combination thereof. For example, monoclonal antibodies can be produced using hybridoma techniques including those known in the art and taught, for example, in Harlow et al., Antibodies: A Laboratory Manual, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988); Hammerling, et al., in: Monoclonal Antibodies and T-Cell Hybridomas 563-681 (Elsevier, N.Y., 1981) (said references incorporated by reference in their entireties). The term "monoclonal antibody" as used herein is not limited to antibodies produced through hybridoma technology. The term "monoclonal antibody" refers to an antibody that is derived from a single clone, including any eukaryotic, prokaryotic, or phage clone, and not the method by which it is produced.

hybridoma technology are routine and well known in the art and are discussed in detail in the Examples. In a non-limiting example, mice can be immunized with a polypeptide of the invention or a cell expressing such peptide. Once an immune response is detected, e.g., antibodies specific for the antigen are detected in the mouse serum, the mouse spleen is harvested and splenocytes isolated. The splenocytes are then fused by well known techniques to any suitable myeloma cells, for example cells from cell line SP20 available from the ATCC. Hybridomas are selected and cloned by limited dilution. The hybridoma clones are then assayed by methods known in the art for cells that secrete antibodies capable of binding a polypeptide of the invention.

Ascites fluid, which generally contains high levels of antibodies, can be generated by immunizing mice with positive hybridoma clones.

[0249] Accordingly, the present invention provides methods of generating monoclonal antibodies as well as antibodies produced by the method comprising culturing a hybridoma cell secreting an antibody of the invention wherein, preferably, the hybridoma is generated by fusing splenocytes isolated from a mouse immunized with an antigen of the invention with myeloma cells and then screening the hybridomas resulting from the fusion for hybridoma clones that secrete an antibody able to bind a polypeptide of the invention.

human B cell lines is transformation using Epstein Barr Virus (EBV). Protocols for generating EBV-transformed B cell lines are commonly known in the art, such as, for example, the protocol outlined in Chapter 7.22 of Current Protocols in Immunology, Coligan et al., Eds., 1994, John Wiley & Sons, NY, which is hereby incorporated in its entirety by reference herein. The source of B cells for transformation is commonly human peripheral blood, but B cells for transformation may also be derived from other sources including, but not limited to, lymph nodes, tonsil, spleen, tumor tissue, and infected tissues. Tissues are generally made into single cell suspensions prior to EBV transformation. Additionally, steps may be taken to either physically remove or inactivate T cells (e.g., by treatment with cyclosporin A) in B cell-containing samples, because T cells from individuals seropositive for anti-EBV antibodies can suppress B cell immortalization by EBV.

In general, the sample containing human B cells is innoculated with EBV, and cultured for 3-4 weeks. A typical source of EBV is the culture supernatant of the B95-8 cell line (ATCC #VR-1492). Physical signs of EBV transformation can generally be seen towards the end of the 3-4 week culture period. By phase-contrast microscopy, transformed cells may appear large, clear, hairy and tend to aggregate in tight clusters of cells. Initially, EBV lines are generally polyclonal. However, over prolonged periods of cell cultures, EBV lines may become monoclonal or polyclonal as a result of the selective outgrowth of particular B cell clones. Alternatively, polyclonal EBV transformed lines may be subcloned (e.g., by limiting dilution culture) or fused with a suitable fusion partner and plated at limiting dilution to obtain

monoclonal B cell lines. Suitable fusion partners for EBV transformed cell lines include mouse myeloma cell lines (e.g., SP2/0, X63-Ag8.653), heteromyeloma cell lines (human x mouse; e.g, SPAM-8, SBC-H20, and CB-F7), and human cell lines (e.g., GM 1500, SKO-007, RPMI 8226, and KR-4). Thus, the present invention also provides a method of generating polyclonal or monoclonal human antibodies against polypeptides of the invention or fragments thereof, comprising EBV-transformation of human B cells.

[0252]Antibody fragments which recognize specific epitopes may be generated by known techniques. For example, Fab and F(ab')2 fragments of the invention may be produced by proteolytic cleavage of immunoglobulin molecules, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')2 fragments). F(ab')2 fragments contain the variable region, the light chain constant region and the CH1 domain of the heavy chain. For example, the antibodies of the present invention can also be generated using various phage display methods known in the art and as discussed in detail in the Examples (e.g., Example 10). In phage display methods, functional antibody domains are displayed on the surface of phage particles which carry the polynucleotide sequences encoding them. In a particular embodiment, such phage can be utilized to display antigen binding domains expressed from a repertoire or combinatorial antibody library (e.g., human or murine). expressing an antigen binding domain that binds the antigen of interest can be selected or identified with antigen, e.g., using labeled antigen or antigen bound or captured to a solid surface or bead. Phage used in these methods are typically filamentous phage including fd and M13 binding domains expressed from phage with Fab, Fv or disulfide stabilized Fv antibody domains recombinantly fused to either the phage gene III or gene VIII protein. Examples of phage display methods that can be used to make the antibodies of the present invention include those disclosed in Brinkman et al., J. Immunol. Methods 182:41-50 (1995); Ames et al., J. Immunol. Methods 184:177-186 (1995); Kettleborough et al., Eur. J. Immunol. 24:952-958 (1994); Persic et al., Gene 187 9-18 (1997); Burton et al., Advances in Immunology 57:191-280 (1994); PCT application No. PCT/GB91/01134; PCT publications WO 90/02809; WO 91/10737; WO 92/01047; WO 92/18619; WO 93/11236; WO 95/15982; WO 95/20401; and U.S. Patent Nos. 5,698,426; 5,223,409; 5,403,484; 5,580,717; 5,427,908; 5,750,753;

5,821,047; 5,571,698; 5,427,908; 5,516,637; 5,780,225; 5,658,727; 5,733,743 and 5,969,108; each of which is incorporated herein by reference in its entirety.

As described in the above references, after phage selection, the antibody coding regions from the phage can be isolated and used to generate whole antibodies, including human antibodies, or any other desired antigen binding fragment, and expressed in any desired host, including mammalian cells, insect cells, plant cells, yeast, and bacteria, e.g., as described in detail below. For example, techniques to recombinantly produce Fab, Fab' and F(ab')2 fragments can also be employed using methods known in the art such as those disclosed in PCT publication WO 92/22324; Mullinax et al., BioTechniques 12(6):864-869 (1992); and Sawai et al., AJRI 34:26-34 (1995); and Better et al., Science 240:1041-1043 (1988) (said references incorporated by reference in their entireties).

[0254] Examples of techniques which can be used to produce single-chain Fvs and antibodies include those described in U.S. Patents 4,946,778 and 5,258,498; Huston et al., Methods in Enzymology 203:46-88 (1991); Shu et al., PNAS 90:7995-7999 (1993); and Skerra et al., Science 240:1038-1040 (1988). For some uses, including in vivo use of antibodies in humans and in vitro detection assays, it may be preferable to use chimeric, humanized, or human antibodies. A chimeric antibody is a molecule in which different portions of the antibody are derived from different animal species, such as antibodies having a variable region derived from a murine monoclonal antibody and a human immunoglobulin constant region. Methods for producing chimeric antibodies are known in the art. See e.g., Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Gillies et al., (1989) J. Immunol. Methods 125:191-202; U.S. Patent Nos. 5,807,715; 4,816,567; and 4,816397, which are incorporated herein by reference in their entirety. Humanized antibodies are antibody molecules from non-human species antibody that binds the desired antigen having one or more complementarity determining regions (CDRs) from the nonhuman species and a framework regions from a human immunoglobulin molecule. Often, framework residues in the human framework regions will be substituted with the corresponding residue from the CDR donor antibody to alter, preferably improve, antigen binding. These framework substitutions are identified by methods well known in the art, e.g., by modeling of the interactions of the CDR and framework

residues to identify framework residues important for antigen binding and sequence comparison to identify unusual framework residues at particular positions. (See, e.g., Queen et al., U.S. Patent No. 5,585,089; Riechmann et al., Nature 332:323 (1988), which are incorporated herein by reference in their entireties.) Antibodies can be humanized using a variety of techniques known in the art including, for example, CDR-grafting (EP 239,400; PCT publication WO 91/09967; U.S. Patent Nos. 5,225,539; 5,530,101; and 5,585,089), veneering or resurfacing (EP 592,106; EP 519,596; Padlan, Molecular Immunology 28(4/5):489-498 (1991); Studnicka et al., Protein Engineering 7(6):805-814 (1994); Roguska. et al., PNAS 91:969-973 (1994)), and chain shuffling (U.S. Patent No. 5,565,332).

[0255] Completely human antibodies are particularly desirable for therapeutic treatment of human patients. Human antibodies can be made by a variety of methods known in the art including phage display methods described above using antibody libraries derived from human immunoglobulin sequences. See also, U.S. Patent Nos. 4,444,887 and 4,716,111; and PCT publications WO 98/46645, WO 98/50433, WO 98/24893, WO 98/16654, WO 96/34096, WO 96/33735, and WO 91/10741; each of which is incorporated herein by reference in its entirety.

Human antibodies can also be produced using transgenic mice which are [0256] incapable of expressing functional endogenous immunoglobulins, but which can express human immunoglobulin genes. For example, the human heavy and light chain immunoglobulin gene complexes may be introduced randomly or by homologous recombination into mouse embryonic stem cells. Alternatively, the human variable region, constant region, and diversity region may be introduced into mouse embryonic stem cells in addition to the human heavy and light chain genes. The mouse heavy and light chain immunoglobulin genes may be rendered non-functional separately or simultaneously with the introduction of human immunoglobulin loci by homologous In particular, homozygous deletion of the JH region prevents recombination. endogenous antibody production. The modified embryonic stem cells are expanded and microinjected into blastocysts to produce chimeric mice. The chimeric mice are then bred to produce homozygous offspring which express human antibodies. The transgenic mice are immunized in the normal fashion with a selected antigen, e.g., all or a portion of a polypeptide of the invention. Monoclonal antibodies directed against

the antigen can be obtained from the immunized, transgenic mice using conventional hybridoma technology. The human immunoglobulin transgenes harbored by the transgenic mice rearrange during B cell differentiation, and subsequently undergo class switching and somatic mutation. Thus, using such a technique, it is possible to produce therapeutically useful IgG, IgA, IgM and IgE antibodies. For an overview of this technology for producing human antibodies, see Lonberg and Huszar, Int. Rev. Immunol. 13:65-93 (1995). For a detailed discussion of this technology for producing human antibodies and human monoclonal antibodies and protocols for producing such antibodies, see, e.g., PCT publications WO 98/24893; WO 92/01047; WO 96/34096; WO 96/33735; European Patent No. 0 598 877; U.S. Patent Nos. 5,413,923; 5,625,126; 5,633,425; 5,569,825; 5,661,016; 5,545,806; 5,814,318; 5,885,793; 5,916,771; 5,939,598; 6,075,181 and 6,114,598, which are incorporated by reference herein in their entirety. In addition, companies such as Abgenix, Inc. (Freemont, CA) and Genpharm (San Jose, CA) can be engaged to provide human antibodies directed against a selected antigen using technology similar to that described above.

[0257] Completely human antibodies which recognize a selected epitope can be generated using a technique referred to as "guided selection." In this approach a selected non-human monoclonal antibody, e.g., a mouse antibody, is used to guide the selection of a completely human antibody recognizing the same epitope. (Jespers et al., Bio/technology 12:899-903 (1988)).

to generate anti-idiotype antibodies that "mimic" polypeptides of the invention using techniques well known to those skilled in the art. (See, e.g., Greenspan & Bona, FASEB J. 7(5):437-444; (1989) and Nissinoff, J. Immunol. 147(8):2429-2438 (1991)). For example, antibodies which bind to and competitively inhibit polypeptide multimerization and/or binding of a polypeptide of the invention to a ligand can be used to generate anti-idiotypes that "mimic" the polypeptide multimerization and/or binding domain and, as a consequence, bind to and neutralize polypeptide and/or its ligand. Such neutralizing anti-idiotypes or Fab fragments of such anti-idiotypes can be used in therapeutic regimens to neutralize polypeptide ligand/receptor. For example, such anti-idiotypic antibodies can be used to bind a polypeptide of the invention and/or to bind its ligand(s)/receptor(s), and thereby block its biological activity.

Alternatively, antibodies which bind to and enhance polypeptide multimerization and/or binding, and/or receptor/ligand multimerization, binding and/or signaling can be used to generate anti-idiotypes that function as agonists of a polypeptide of the invention and/or its ligand/receptor. Such agonistic anti-idiotypes or Fab fragments of such anti-idiotypes can be used in therapeutic regimens as agonists of the polypeptides of the invention or its ligand(s)/receptor(s). For example, such anti-idiotypic antibodies can be used to bind a polypeptide of the invention and/or to bind its ligand(s)/receptor(s), and thereby promote or enhance its biological activity.

Intrabodies of the invention can be produced using methods known in the art, such as those disclosed and reviewed in Chen et al., Hum. Gene Ther. 5:595-601 (1994); Marasco, W.A., Gene Ther. 4:11-15 (1997); Rondon and Marasco, Annu. Rev. Microbiol. 51:257-283 (1997); Proba et al., J. Mol. Biol. 275:245-253 (1998); Cohen et al., Oncogene 17:2445-2456 (1998); Ohage and Steipe, J. Mol. Biol. 291:1119-1128 (1999); Ohage et al., J. Mol. Biol. 291:1129-1134 (1999); Wirtz and Steipe, Protein Sci. 8:2245-2250 (1999); Zhu et al., J. Immunol. Methods 231:207-222 (1999); and references cited therein.

Polynucleotides Encoding Antibodies

The invention further provides polynucleotides comprising a nucleotide sequence encoding an antibody of the invention and fragments thereof. The invention also encompasses polynucleotides that hybridize under stringent or alternatively, under lower stringency hybridization conditions, e.g., as defined *supra*, to polynucleotides that encode an antibody, preferably, that specifically binds to a polypeptide of the invention, preferably, an antibody that binds to a polypeptide having the amino acid sequence of SEQ ID NO:Y, to a polypeptide encoded by a portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or to a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0261] The polynucleotides may be obtained, and the nucleotide sequence of the polynucleotides determined, by any method known in the art. For example, if the nucleotide sequence of the antibody is known, a polynucleotide encoding the antibody may be assembled from chemically synthesized oligonucleotides (e.g., as described in Kutmeier et al., BioTechniques 17:242 (1994)), which, briefly, involves the synthesis

of overlapping oligonucleotides containing portions of the sequence encoding the antibody, annealing and ligating of those oligonucleotides, and then amplification of the ligated oligonucleotides by PCR.

In a close containing an antibody may be generated from nucleic acid from a suitable source. If a clone containing a nucleic acid encoding a particular antibody is not available, but the sequence of the antibody molecule is known, a nucleic acid encoding the immunoglobulin may be chemically synthesized or obtained from a suitable source (e.g., an antibody cDNA library, or a cDNA library generated from, or nucleic acid, preferably poly A+RNA, isolated from, any tissue or cells expressing the antibody, such as hybridoma cells selected to express an antibody of the invention) by PCR amplification using synthetic primers hybridizable to the 3' and 5' ends of the sequence or by cloning using an oligonucleotide probe specific for the particular gene sequence to identify, e.g., a cDNA clone from a cDNA library that encodes the antibody. Amplified nucleic acids generated by PCR may then be cloned into replicable cloning vectors using any method well known in the art.

the antibody is determined, the nucleotide sequence of the antibody may be manipulated using methods well known in the art for the manipulation of nucleotide sequences, e.g., recombinant DNA techniques, site directed mutagenesis, PCR, etc. (see, for example, the techniques described in Sambrook et al., 1990, Molecular Cloning, A Laboratory Manual, 2d Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY and Ausubel et al., eds., 1998, Current Protocols in Molecular Biology, John Wiley & Sons, NY, which are both incorporated by reference herein in their entireties), to generate antibodies having a different amino acid sequence, for example to create amino acid substitutions, deletions, and/or insertions.

[0264] In a specific embodiment, the amino acid sequence of the heavy and/or light chain variable domains may be inspected to identify the sequences of the complementarity determining regions (CDRs) by methods that are well know in the art, e.g., by comparison to known amino acid sequences of other heavy and light chain variable regions to determine the regions of sequence hypervariability. Using routine recombinant DNA techniques, one or more of the CDRs may be inserted within framework regions, e.g., into human framework regions to humanize a non-human

antibody, as described *supra*. The framework regions may be naturally occurring or consensus framework regions, and preferably human framework regions (see, e.g., Chothia et al., J. Mol. Biol. 278: 457-479 (1998) for a listing of human framework regions). Preferably, the polynucleotide generated by the combination of the framework regions and CDRs encodes an antibody that specifically binds a polypeptide of the invention. Preferably, as discussed *supra*, one or more amino acid substitutions may be made within the framework regions, and, preferably, the amino acid substitutions improve binding of the antibody to its antigen. Additionally, such methods may be used to make amino acid substitutions or deletions of one or more variable region cysteine residues participating in an intrachain disulfide bond to generate antibody molecules lacking one or more intrachain disulfide bonds. Other alterations to the polynucleotide are encompassed by the present invention and within the skill of the art.

[0265] In addition, techniques developed for the production of "chimeric antibodies" (Morrison et al., Proc. Natl. Acad. Sci. 81:851-855 (1984); Neuberger et al., Nature 312:604-608 (1984); Takeda et al., Nature 314:452-454 (1985)) by splicing genes from a mouse antibody molecule of appropriate antigen specificity together with genes from a human antibody molecule of appropriate biological activity can be used. As described *supra*, a chimeric antibody is a molecule in which different portions are derived from different animal species, such as those having a variable region derived from a murine mAb and a human immunoglobulin constant region, e.g., humanized antibodies.

Alternatively, techniques described for the production of single chain antibodies (U.S. Patent No. 4,946,778; Bird, Science 242:423- 42 (1988); Huston et al., Proc. Natl. Acad. Sci. USA 85:5879-5883 (1988); and Ward et al., Nature 334:544-54 (1989)) can be adapted to produce single chain antibodies. Single chain antibodies are formed by linking the heavy and light chain fragments of the Fv region via an amino acid bridge, resulting in a single chain polypeptide. Techniques for the assembly of functional Fv fragments in *E. coli* may also be used (Skerra et al., Science 242:1038-1041 (1988)).

The antibodies of the invention can be produced by any method known in the art for the synthesis of antibodies, in particular, by chemical synthesis or preferably, by recombinant expression techniques. Methods of producing antibodies include, but are not limited to, hybridoma technology, EBV transformation, and other methods discussed herein as well as through the use recombinant DNA technology, as discussed below.

[0268] Recombinant expression of an antibody of the invention, or fragment, derivative or analog thereof, (e.g., a heavy or light chain of an antibody of the invention or a single chain antibody of the invention), requires construction of an expression vector containing a polynucleotide that encodes the antibody. Once a polynucleotide encoding an antibody molecule or a heavy or light chain of an antibody, or portion thereof (preferably containing the heavy or light chain variable domain), of the invention has been obtained, the vector for the production of the antibody molecule may be produced by recombinant DNA technology using techniques well known in the art. Thus, methods for preparing a protein by expressing a polynucleotide containing an antibody encoding nucleotide sequence are described herein. Methods which are well known to those skilled in the art can be used to construct expression vectors containing antibody coding sequences and appropriate transcriptional and translational control signals. These methods include, for example, in vitro recombinant DNA techniques, synthetic techniques, and in vivo genetic The invention, thus, provides replicable vectors comprising a recombination. nucleotide sequence encoding an antibody molecule of the invention, or a heavy or light chain thereof, or a heavy or light chain variable domain, operably linked to a promoter. Such vectors may include the nucleotide sequence encoding the constant region of the antibody molecule (see, e.g., PCT Publication WO 86/05807; PCT Publication WO 89/01036; and U.S. Patent No. 5,122,464) and the variable domain of the antibody may be cloned into such a vector for expression of the entire heavy or light chain.

[0269] The expression vector is transferred to a host cell by conventional techniques and the transfected cells are then cultured by conventional techniques to produce an antibody of the invention. Thus, the invention includes host cells containing a polynucleotide encoding an antibody of the invention, or a heavy or light

chain thereof, or a single chain antibody of the invention, operably linked to a heterologous promoter. In preferred embodiments for the expression of double-chained antibodies, vectors encoding both the heavy and light chains may be co-expressed in the host cell for expression of the entire immunoglobulin molecule, as detailed below.

[0270] A variety of host-expression vector systems may be utilized to express the antibody molecules of the invention. Such host-expression systems represent vehicles by which the coding sequences of interest may be produced and subsequently purified, but also represent cells which may, when transformed or transfected with the appropriate nucleotide coding sequences, express an antibody molecule of the invention in situ. These include but are not limited to microorganisms such as bacteria (e.g., E. coli, B. subtilis) transformed with recombinant bacteriophage DNA, plasmid DNA or cosmid DNA expression vectors containing antibody coding sequences; yeast (e.g., Saccharomyces, Pichia) transformed with recombinant yeast expression vectors containing antibody coding sequences; insect cell systems infected with recombinant virus expression vectors (e.g., baculovirus) containing antibody coding sequences; plant cell systems infected with recombinant virus expression vectors (e.g., cauliflower mosaic virus, CaMV; tobacco mosaic virus, TMV) or transformed with recombinant plasmid expression vectors (e.g., Ti plasmid) containing antibody coding sequences; or mammalian cell systems (e.g., COS, CHO, BHK, 293, 3T3 cells) harboring recombinant expression constructs containing promoters derived from the genome of mammalian cells (e.g., metallothionein promoter) or from mammalian viruses (e.g., the adenovirus late promoter; the vaccinia virus 7.5K promoter). Preferably, bacterial cells such as Escherichia coli, and more preferably, eukaryotic cells, especially for the expression of whole recombinant antibody molecule, are used for the expression of a recombinant antibody molecule. For example, mammalian cells such as Chinese hamster ovary cells (CHO), in conjunction with a vector such as the major intermediate early gene promoter element from human cytomegalovirus is an effective expression system for antibodies (Foecking et al., Gene 45:101 (1986); Cockett et al., Bio/Technology 8:2 (1990)).

[0271] In bacterial systems, a number of expression vectors may be advantageously selected depending upon the use intended for the antibody molecule

being expressed. For example, when a large quantity of such a protein is to be produced, for the generation of pharmaceutical compositions of an antibody molecule, vectors which direct the expression of high levels of fusion protein products that are readily purified may be desirable. Such vectors include, but are not limited, to the E. coli expression vector pUR278 (Ruther et al., EMBO J. 2:1791 (1983)), in which the antibody coding sequence may be ligated individually into the vector in frame with the lac Z coding region so that a fusion protein is produced; pIN vectors (Inouye & Inouye, Nucleic Acids Res. 13:3101-3109 (1985); Van Heeke & Schuster, J. Biol. Chem. 24:5503-5509 (1989)); and the like. pGEX vectors may also be used to express foreign polypeptides as fusion proteins with glutathione S-transferase (GST). In general, such fusion proteins are soluble and can easily be purified from lysed cells by adsorption and binding to matrix glutathione-agarose beads followed by elution in the presence of free glutathione. The pGEX vectors are designed to include thrombin or factor Xa protease cleavage sites so that the cloned target gene product can be released from the GST moiety.

[0272] In an insect system, Autographa californica nuclear polyhedrosis virus (AcNPV) is used as a vector to express foreign genes. The virus grows in Spodoptera frugiperda cells. The antibody coding sequence may be cloned individually into non-essential regions (for example the polyhedrin gene) of the virus and placed under control of an AcNPV promoter (for example the polyhedrin promoter).

be utilized. In cases where an adenovirus is used as an expression vector, the antibody coding sequence of interest may be ligated to an adenovirus transcription/translation control complex, e.g., the late promoter and tripartite leader sequence. This chimeric gene may then be inserted in the adenovirus genome by in vitro or in vivo recombination. Insertion in a non- essential region of the viral genome (e.g., region E1 or E3) will result in a recombinant virus that is viable and capable of expressing the antibody molecule in infected hosts. (e.g., see Logan & Shenk, Proc. Natl. Acad. Sci. USA 81:355-359 (1984)). Specific initiation signals may also be required for efficient translation of inserted antibody coding sequences. These signals include the ATG initiation codon and adjacent sequences. Furthermore, the initiation codon must be in phase with the reading frame of the desired coding sequence to

ensure translation of the entire insert. These exogenous translational control signals and initiation codons can be of a variety of origins, both natural and synthetic. The efficiency of expression may be enhanced by the inclusion of appropriate transcription enhancer elements, transcription terminators, etc. (see Bittner et al., Methods in Enzymol. 153:51-544 (1987)).

[0274] In addition, a host cell strain may be chosen which modulates the expression of the inserted sequences, or modifies and processes the gene product in the specific fashion desired. Such modifications (e.g., glycosylation) and processing (e.g., cleavage) of protein products may be important for the function of the protein. Different host cells have characteristic and specific mechanisms for the post-translational processing and modification of proteins and gene products. Appropriate cell lines or host systems can be chosen to ensure the correct modification and processing of the foreign protein expressed. To this end, eukaryotic host cells which possess the cellular machinery for proper processing of the primary transcript, glycosylation, and phosphorylation of the gene product may be used. Such mammalian host cells include but are not limited to CHO, VERY, BHK, Hela, COS, MDCK, 293, 3T3, WI38, and in particular, breast cancer cell lines such as, for example, BT483, Hs578T, HTB2, BT20 and T47D, and normal mammary gland cell line such as, for example, CRL7030 and Hs578Bst.

For long-term, high-yield production of recombinant proteins, stable expression is preferred. For example, cell lines which stably express the antibody molecule may be engineered. Rather than using expression vectors which contain viral origins of replication, host cells can be transformed with DNA controlled by appropriate expression control elements (e.g., promoter, enhancer, sequences, transcription terminators, polyadenylation sites, etc.), and a selectable marker. Following the introduction of the foreign DNA, engineered cells may be allowed to grow for 1-2 days in an enriched media, and then are switched to a selective media. The selectable marker in the recombinant plasmid confers resistance to the selection and allows cells to stably integrate the plasmid into their chromosomes and grow to form foci which in turn can be cloned and expanded into cell lines. This method may advantageously be used to engineer cell lines which express the antibody molecule.

Such engineered cell lines may be particularly useful in screening and evaluation of compounds that interact directly or indirectly with the antibody molecule.

A number of selection systems may be used, including but not limited to [0276] the herpes simplex virus thymidine kinase (Wigler et al., Cell 11:223 (1977)), hypoxanthine-guanine phosphoribosyltransferase (Szybalska & Szybalski, Proc. Natl. Acad. Sci. USA 48:202 (1992)), and adenine phosphoribosyltransferase (Lowy et al., Cell 22:817 (1980)) genes can be employed in tk-, hgprt- or aprt- cells, respectively. Also, antimetabolite resistance can be used as the basis of selection for the following genes: dhfr, which confers resistance to methotrexate (Wigler et al., Natl. Acad. Sci. USA 77:357 (1980); O'Hare et al., Proc. Natl. Acad. Sci. USA 78:1527 (1981)); gpt, which confers resistance to mycophenolic acid (Mulligan & Berg, Proc. Natl. Acad. Sci. USA 78:2072 (1981)); neo, which confers resistance to the aminoglycoside G-418 Clinical Pharmacy 12:488-505; Wu and Wu, Biotherapy 3:87-95 (1991); Tolstoshev, Ann. Rev. Pharmacol. Toxicol. 32:573-596 (1993); Mulligan, Science 260:926-932 (1993); and Morgan and Anderson, Ann. Rev. Biochem. 62:191-217 (1993); TIB TECH 11(5):155-215 (1993)); and hygro, which confers resistance to hygromycin (Santerre et al., Gene 30:147 (1984)). Methods commonly known in the art of recombinant DNA technology may be routinely applied to select the desired recombinant clone, and such methods are described, for example, in Ausubel et al. (eds.), Current Protocols in Molecular Biology, John Wiley & Sons, NY (1993); Kriegler, Gene Transfer and Expression, A Laboratory Manual, Stockton Press, NY (1990); and in Chapters 12 and 13, Dracopoli et al. (eds), Current Protocols in Human Genetics, John Wiley & Sons, NY (1994); Colberre-Garapin et al., J. Mol. Biol. 150:1 (1981), which are incorporated by reference herein in their entireties.

[0277] The expression levels of an antibody molecule can be increased by vector amplification (for a review, see Bebbington and Hentschel, The use of vectors based on gene amplification for the expression of cloned genes in mammalian cells in DNA cloning, Vol.3. (Academic Press, New York, 1987)). When a marker in the vector system expressing antibody is amplifiable, increase in the level of inhibitor present in culture of host cell will increase the number of copies of the marker gene. Since the amplified region is associated with the antibody gene, production of the antibody will also increase (Crouse et al., Mol. Cell. Biol. 3:257 (1983)).

Vectors which use glutamine synthase (GS) or DHFR as the selectable [0278] markers can be amplified in the presence of the drugs methionine sulphoximine or methotrexate, respectively. An advantage of glutamine synthase based vectors are the availabilty of cell lines (e.g., the murine myeloma cell line, NS0) which are glutamine synthase negative. Glutamine synthase expression systems can also function in glutamine synthase expressing cells (e.g., Chinese Hamster Ovary (CHO) cells) by providing additional inhibitor to prevent the functioning of the endogenous gene. A glutamine synthase expression system and components thereof are detailed in PCT publications: WO87/04462; WO86/05807; WO89/01036; WO89/10404; and WO91/06657 which are incorporated in their entireties by reference herein. Additionally, glutamine synthase expression vectors that may be used according to the present invention are commercially available from suplliers, including, for example Lonza Biologics, Inc. (Portsmouth, NH). Expression and production of monoclonal antibodies using a GS expression system in murine myeloma cells is described in Bebbington et al., Bio/technology 10:169(1992) and in Biblia and Robinson Biotechnol. Prog. 11:1 (1995) which are incorporated in their entirities by reference herein.

[0279] The host cell may be co-transfected with two expression vectors of the invention, the first vector encoding a heavy chain derived polypeptide and the second vector encoding a light chain derived polypeptide. The two vectors may contain identical selectable markers which enable equal expression of heavy and light chain polypeptides. Alternatively, a single vector may be used which encodes, and is capable of expressing, both heavy and light chain polypeptides. In such situations, the light chain should be placed before the heavy chain to avoid an excess of toxic free heavy chain (Proudfoot, Nature 322:52 (1986); Kohler, Proc. Natl. Acad. Sci. USA 77:2197 (1980)). The coding sequences for the heavy and light chains may comprise cDNA or genomic DNA.

[0280] Once an antibody molecule of the invention has been produced by an animal, chemically synthesized, or recombinantly expressed, it may be purified by any method known in the art for purification of an immunoglobulin molecule, for example, by chromatography (e.g., ion exchange, affinity, particularly by affinity for the specific antigen after Protein A, and sizing column chromatography), centrifugation,

differential solubility, or by any other standard technique for the purification of proteins. In addition, the antibodies of the present invention or fragments thereof can be fused to heterologous polypeptide sequences described herein or otherwise known in the art, to facilitate purification.

The present invention encompasses antibodies recombinantly fused or [0281] chemically conjugated (including both covalently and non-covalently conjugations) to a polypeptide (or portion thereof, preferably at least 10, 20, 30, 40, 50, 60, 70, 80, 90 or 100 amino acids of the polypeptide) of the present invention to generate fusion proteins. The fusion does not necessarily need to be direct, but may occur through linker sequences. The antibodies may be specific for antigens other than polypeptides (or portion thereof, preferably at least 10, 20, 30, 40, 50, 60, 70, 80, 90 or 100 amino acids of the polypeptide) of the present invention. For example, antibodies may be used to target the polypeptides of the present invention to particular cell types, either in vitro or in vivo, by fusing or conjugating the polypeptides of the present invention to antibodies specific for particular cell surface receptors. Antibodies fused or conjugated to the polypeptides of the present invention may also be used in in vitro immunoassays and purification methods using methods known in the art. See e.g., Harbor et al., supra, and PCT publication WO 93/21232; EP 439,095; Naramura et al., Immunol. Lett. 39:91-99 (1994); U.S. Patent 5,474,981; Gillies et al., PNAS 89:1428-1432 (1992); Fell et al., J. Immunol. 146:2446-2452 (1991), which are incorporated by reference in their entireties.

[0282] The present invention further includes compositions comprising the polypeptides of the present invention fused or conjugated to antibody domains other than the variable regions. For example, the polypeptides of the present invention may be fused or conjugated to an antibody Fc region, or portion thereof. The antibody portion fused to a polypeptide of the present invention may comprise the constant region, hinge region, CH1 domain, CH2 domain, and CH3 domain or any combination of whole domains or portions thereof. The polypeptides may also be fused or conjugated to the above antibody portions to form multimers. For example, Fc portions fused to the polypeptides of the present invention can form dimers through disulfide bonding between the Fc portions. Higher multimeric forms can be made by fusing the polypeptides to portions of IgA and IgM. Methods for fusing or

conjugating the polypeptides of the present invention to antibody portions are known in the art. See, e.g., U.S. Patent Nos. 5,336,603; 5,622,929; 5,359,046; 5,349,053; 5,447,851; 5,112,946; EP 307,434; EP 367,166; PCT publications WO 96/04388; WO 91/06570; Ashkenazi et al., Proc. Natl. Acad. Sci. USA 88:10535-10539 (1991); Zheng et al., J. Immunol. 154:5590-5600 (1995); and Vil et al., Proc. Natl. Acad. Sci. USA 89:11337- 11341 (1992) (said references incorporated by reference in their entireties).

[0283] As discussed, supra, the polypeptides corresponding to a polypeptide, polypeptide fragment, or a variant of SEQ ID NO:Y may be fused or conjugated to the above antibody portions to increase the in vivo half life of the polypeptides or for use in immunoassays using methods known in the art. Further, the polypeptides corresponding to SEQ ID NO:Y may be fused or conjugated to the above antibody portions to facilitate purification. One reported example describes chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. See EP 394,827; Traunecker et al., Nature 331:84-86 (1988). The polypeptides of the present invention fused or conjugated to an antibody having disulfide- linked dimeric structures (due to the IgG) may also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. See, for example, Fountoulakis et al., J. Biochem. 270:3958-3964 (1995). In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties. See, for example, EP A 232,262. Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, Bennett et al., J. Molecular Recognition 8:52-58 (1995); Johanson et al., J. Biol. Chem. 270:9459-9471 (1995)).

[0284] Moreover, the antibodies or fragments thereof of the present invention can be fused to marker sequences, such as a peptide to facilitate purification. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the

tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexahistidine provides for convenient purification of the fusion protein. Other peptide tags useful for purification include, but are not limited to, the "HA" tag, which corresponds to an epitope derived from the influenza hemagglutinin protein (Wilson et al., Cell 37:767 (1984)) and the "flag" tag.

[0285] The present invention further encompasses antibodies or fragments thereof conjugated to a diagnostic or therapeutic agent. The antibodies can diagnostically to, for example, monitor the development or progression of a tumor as part of a clinical testing procedure to, e.g., determine the efficacy of a given treatment regimen. Detection can be facilitated by coupling the antibody to a detectable substance. Examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, radioactive materials, positron emitting metals using various positron emission tomographies, and nonradioactive paramagnetic metal ions. The detectable substance may be coupled or conjugated either directly to the antibody (or fragment thereof) or indirectly, through an intermediate (such as, for example, a linker known in the art) using techniques known in the art. See, for example, U.S. Patent No. 4,741,900 for metal ions which can be conjugated to antibodies for use as diagnostics according to the present invention.

therapeutic moiety such as a cytotoxin, e.g., a cytostatic or cytocidal agent, a therapeutic agent or a radioactive metal ion, e.g., alpha-emitters such as, for example, 213Bi. A cytotoxin or cytotoxic agent includes any agent that is detrimental to cells. Examples include paclitaxol, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, tenoposide, vincristine, vinblastine, colchicin, doxorubicin, daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone, glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs or homologs thereof. Therapeutic agents include, but are not limited to, antimetabolites (e.g., methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating

agents (e.g., mechlorethamine, thioepa chlorambucil, melphalan, carmustine (BSNU) and lomustine (CCNU), cyclothosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cis- dichlorodiamine platinum (II) (DDP) cisplatin), anthracyclines (e.g., daunorubicin (formerly daunomycin) and doxorubicin), antibiotics (e.g., dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)), and anti-mitotic agents (e.g., vincristine and vinblastine).

[0287] The conjugates of the invention can be used for modifying a given biological response, the therapeutic agent or drug moiety is not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological activity. Such proteins may include, for example, a toxin such as abrin, ricin A, pseudomonas exotoxin, or diphtheria toxin; a protein such as tumor necrosis factor, a-interferon, B-interferon, nerve growth factor, platelet derived growth factor, tissue plasminogen activator, an apoptotic agent, e.g., TNF-alpha, TNF-beta, AIM I (See, International Publication No. WO 97/33899), AIM II (See, International Publication No. WO 97/34911), Fas Ligand (Takahashi et al., Int. Immunol., 6:1567-1574 (1994)), VEGI (See, International Publication No. WO 99/23105), a thrombotic agent or an anti- angiogenic agent, e.g., angiostatin or endostatin; or, biological response modifiers such as, for example, lymphokines, interleukin-1 ("IL-1"), interleukin-2 ("IL-2"), interleukin-6 ("IL-6"), granulocyte macrophage colony stimulating factor ("GM-CSF"), granulocyte colony stimulating factor ("G-CSF"), or other growth factors.

[0288] Antibodies may also be attached to solid supports, which are particularly useful for immunoassays or purification of the target antigen. Such solid supports include, but are not limited to, glass, cellulose, polyacrylamide, nylon, polystyrene, polyvinyl chloride or polypropylene.

known. See, for example., Arnon et al., "Monoclonal Antibodies For Immunotargeting Of Drugs In Cancer Therapy", in Monoclonal Antibodies And Cancer Therapy, Reisfeld et al. (eds.), pp. 243-56 (Alan R. Liss, Inc. 1985); Hellstrom et al., "Antibodies For Drug Delivery", in Controlled Drug Delivery (2nd Ed.), Robinson et al. (eds.), pp. 623-53 (Marcel Dekker, Inc. 1987); Thorpe, "Antibody Carriers Of Cytotoxic Agents In Cancer Therapy: A Review", in Monoclonal

Antibodies '84: Biological And Clinical Applications, Pinchera et al. (eds.), pp. 475-506 (1985); "Analysis, Results, And Future Prospective Of The Therapeutic Use Of Radiolabeled Antibody In Cancer Therapy", in Monoclonal Antibodies For Cancer Detection And Therapy, Baldwin et al. (eds.), pp. 303-16 (Academic Press 1985), and Thorpe et al., "The Preparation And Cytotoxic Properties Of Antibody-Toxin Conjugates", Immunol. Rev. 62:119-58 (1982).

[0290] Alternatively, an antibody can be conjugated to a second antibody to form an antibody heteroconjugate as described by Segal in U.S. Patent No. 4,676,980, which is incorporated herein by reference in its entirety.

[0291] An antibody, with or without a therapeutic moiety conjugated to it, administered alone or in combination with cytotoxic factor(s) and/or cytokine(s) can be used as a therapeutic.

Immunophenotyping

The antibodies of the invention may be utilized for immunophenotyping of cell lines and biological samples. Translation products of the genes of the present invention may be useful as cell specific markers, or more specifically as cellular markers that are differentially expressed at various stages of differentiation and/or maturation of particular cell types. Monoclonal antibodies directed against a specific epitope, or combination of epitopes, will allow for the screening of cellular populations expressing the marker. Various techniques can be utilized using monoclonal antibodies to screen for cellular populations expressing the marker(s), and include magnetic separation using antibody-coated magnetic beads, "panning" with antibody attached to a solid matrix (i.e., plate), and flow cytometry (See, e.g., U.S. Patent 5,985,660; and Morrison et al., Cell, 96:737-49 (1999)).

[0293] These techniques allow for the screening of particular populations of cells, such as might be found with hematological malignancies (i.e. minimal residual disease (MRD) in acute leukemic patients) and "non-self" cells in transplantations to prevent Graft-versus-Host Disease (GVHD). Alternatively, these techniques allow for the screening of hematopoietic stem and progenitor cells capable of undergoing proliferation and/or differentiation, as might be found in human umbilical cord blood.

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nylon, blocking the membrane in blocking solution (e.g., PBS with 3% BSA or non-fat milk), washing the membrane in washing buffer (e.g., PBS-Tween 20), blocking the membrane with primary antibody (the antibody of interest) diluted in blocking buffer, washing the membrane in washing buffer, blocking the membrane with a secondary antibody (which recognizes the primary antibody, e.g., an anti-human antibody) conjugated to an enzymatic substrate (e.g., horseradish peroxidase or alkaline phosphatase) or radioactive molecule (e.g., 32P or 125I) diluted in blocking buffer, washing the membrane in wash buffer, and detecting the presence of the antigen. One of skill in the art would be knowledgeable as to the parameters that can be modified to increase the signal detected and to reduce the background noise. For further discussion regarding western blot protocols see, e.g., Ausubel et al., eds., (1994), Current Protocols in Molecular Biology, Vol. 1, John Wiley & Sons, Inc., New York, section 10.8.1.

[0297] ELISAs comprise preparing antigen, coating the well of a 96 well microtiter plate with the antigen, adding the antibody of interest conjugated to a detectable compound such as an enzymatic substrate (e.g., horseradish peroxidase or alkaline phosphatase) to the well and incubating for a period of time, and detecting the presence of the antigen. In ELISAs the antibody of interest does not have to be conjugated to a detectable compound; instead, a second antibody (which recognizes the antibody of interest) conjugated to a detectable compound may be added to the well. Further, instead of coating the well with the antigen, the antibody may be coated to the well. In this case, a second antibody conjugated to a detectable compound may be added following the addition of the antigen of interest to the coated well. One of skill in the art would be knowledgeable as to the parameters that can be modified to increase the signal detected as well as other variations of ELISAs known in the art. For further discussion regarding ELISAs see, e.g., Ausubel et al., eds., (1994), Current Protocols in Molecular Biology, Vol. 1, John Wiley & Sons, Inc., New York, section 11.2.1.

[0298] The binding affinity of an antibody to an antigen and the off-rate of an antibody-antigen interaction can be determined by competitive binding assays. One example of a competitive binding assay is a radioimmunoassay comprising the incubation of labeled antigen (e.g., 3H or 125I) with the antibody of interest in the

presence of increasing amounts of unlabeled antigen, and the detection of the antibody bound to the labeled antigen. The affinity of the antibody of interest for a particular antigen and the binding off-rates can be determined from the data by scatchard plot analysis. Competition with a second antibody can also be determined using radioimmunoassays. In this case, the antigen is incubated with antibody of interest conjugated to a labeled compound (e.g., 3H or 125I) in the presence of increasing amounts of an unlabeled second antibody.

[0299] Antibodies of the invention may be characterized using immunocytochemisty methods on cells (e.g., mammalian cells, such as CHO cells) transfected with a vector enabling the expression of a reproductive system antigen or with vector alone using techniques commonly known in the art. Antibodies that bind reproductive system antigen transfected cells, but not vector-only transfected cells, are reproductive system antigen specific.

Therapeutic Uses

The present invention is further directed to antibody-based therapies which [0300] involve administering antibodies of the invention to an animal, preferably a mammal, and most preferably a human, patient for treating one or more of the disclosed diseases, disorders, or conditions. Therapeutic compounds of the invention include, but are not limited to, antibodies of the invention (including fragments, analogs and derivatives thereof as described herein) and nucleic acids encoding antibodies of the invention (including fragments, analogs and derivatives thereof and anti-idiotypic antibodies as described herein). The antibodies of the invention can be used to treat, inhibit or prevent diseases, disorders or conditions associated with aberrant expression and/or activity of a polypeptide of the invention, including, but not limited to, any one or more of the diseases, disorders, or conditions described herein. The treatment and/or prevention of diseases, disorders, or conditions associated with aberrant expression and/or activity of a polypeptide of the invention includes, but is not limited to, alleviating symptoms associated with those diseases, disorders or conditions. Antibodies of the invention may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

[0301] In a specific and preferred embodiment, the present invention is directed to antibody-based therapies which involve administering antibodies of the invention to an animal, preferably a mammal, and most preferably a human, patient for treating one or more of the diseases, disorders, or conditions of the reproductive system, including, but not limited to, injuries and trauma, infections, neoplastic disorders, congenital defects, and diseases or disorders which result in infertility, complications with pregnancy, labor, or parturition, postpartum difficulties, and as listed below in the section entitled "Reproductive System Disorders". Therapeutic compounds of the invention include, but are not limited to, antibodies of the invention (e.g., antibodies directed to the full length protein expressed on the cell surface of a mammalian cell; antibodies directed to an epitope of a reproductive system associated polypeptide of the invention (such as, a linear epitope (shown in Table 1A, column 6) or a conformational epitope), including fragments, analogs and derivatives thereof as described herein) and nucleic acids encoding antibodies of the invention (including fragments, analogs and derivatives thereof and anti-idiotypic antibodies as described herein). The antibodies of the invention can be used to treat, inhibit or prevent diseases, disorders or conditions associated with aberrant expression and/or activity of a polypeptide of the invention, including, but not limited to, any one or more of the diseases, disorders, or conditions of the reproductive system described herein. The treatment and/or prevention of diseases, disorders, or conditions of the reproductive system associated with aberrant expression and/or activity of a polypeptide of the invention includes, but is not limited to, alleviating symptoms associated with those diseases, disorders or conditions. Antibodies of the invention may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

[0302] A summary of the ways in which the antibodies of the present invention may be used therapeutically includes binding polynucleotides or polypeptides of the present invention locally or systemically in the body or by direct cytotoxicity of the antibody, e.g. as mediated by complement (CDC) or by effector cells (ADCC). Some of these approaches are described in more detail below. Armed with the teachings provided herein, one of ordinary skill in the art will know how to use the antibodies of the present invention for diagnostic, monitoring or therapeutic purposes without undue experimentation.

[0303] The antibodies of this invention may be advantageously utilized in combination with other monoclonal or chimeric antibodies, or with lymphokines or hematopoietic growth factors (such as, e.g., IL-2, IL-3 and IL-7), for example, which serve to increase the number or activity of effector cells which interact with the antibodies.

[0304] The antibodies of the invention may be administered alone or in combination with other types of treatments (e.g., radiation therapy, chemotherapy, hormonal therapy, immunotherapy and anti-tumor agents). Generally, administration of products of a species origin or species reactivity (in the case of antibodies) that is the same species as that of the patient is preferred. Thus, in a preferred embodiment, human antibodies, fragments derivatives, analogs, or nucleic acids, are administered to a human patient for therapy or prophylaxis.

It is preferred to use high affinity and/or potent *in vivo* inhibiting and/or neutralizing antibodies against polypeptides or polynucleotides of the present invention, fragments or regions thereof, for both immunoassays directed to and therapy of disorders related to polynucleotides or polypeptides, including fragments thereof, of the present invention. Such antibodies, fragments, or regions, will preferably have an affinity for polynucleotides or polypeptides of the invention, including fragments thereof. Preferred binding affinities include those with a dissociation constant or Kd less than 5 X 10⁻² M, 10⁻² M, 5 X 10⁻³ M, 10⁻³ M, 5 X 10⁻⁴ M, 10⁻⁴ M, 5 X 10⁻⁵ M, 10⁻⁵ M, 5 X 10⁻⁶ M, 10⁻⁶ M, 5 X 10⁻¹⁷ M, 10⁻⁷ M, 5 X 10⁻⁸ M, 10⁻⁸ M, 5 X 10⁻⁹ M, 10⁻⁹ M, 5 X 10⁻¹⁰ M, 10⁻¹⁰ M, 5 X 10⁻¹¹ M, 10⁻¹¹ M, 5 X 10⁻¹² M, 10⁻¹³ M, 5 X 10⁻¹³ M, 10⁻¹³ M, 5 X 10⁻¹⁴ M, 10⁻¹⁴ M, 5 X 10⁻¹⁵ M, and 10⁻¹⁵ M.

Gene Therapy

[0306] In a specific embodiment, nucleic acids comprising sequences encoding antibodies or functional derivatives thereof, are administered to treat, inhibit or prevent a disease or disorder associated with aberrant expression and/or activity of a polypeptide of the invention, by way of gene therapy. Gene therapy refers to therapy performed by the administration to a subject of an expressed or expressible nucleic acid. In this embodiment of the invention, the nucleic acids produce their encoded protein that mediates a therapeutic effect.

[0307] Any of the methods for gene therapy available in the art can be used according to the present invention. Exemplary methods are described below.

[0308] For general reviews of the methods of gene therapy, see Goldspiel et al., Clinical Pharmacy 12:488-505 (1993); Wu and Wu, Biotherapy 3:87-95 (1991); Tolstoshev, Ann. Rev. Pharmacol. Toxicol. 32:573-596 (1993); Mulligan, Science 260:926-932 (1993); and Morgan and Anderson, Ann. Rev. Biochem. 62:191-217 (1993); May, TIBTECH 11(5):155-215 (1993). Methods commonly known in the art of recombinant DNA technology which can be used are described in Ausubel et al. (eds.), Current Protocols in Molecular Biology, John Wiley & Sons, NY (1993); and Kriegler, Gene Transfer and Expression, A Laboratory Manual, Stockton Press, NY (1990).

[0309] In a preferred embodiment, the compound comprises nucleic acid sequences encoding an antibody, said nucleic acid sequences being part of expression vectors that express the antibody or fragments or chimeric proteins or heavy or light chains thereof in a suitable host. In particular, such nucleic acid sequences have promoters operably linked to the antibody coding region, said promoter being inducible or constitutive, and, optionally, tissue-specific. In another particular embodiment, nucleic acid molecules are used in which the antibody coding sequences and any other desired sequences are flanked by regions that promote homologous recombination at a desired site in the genome, thus providing for intrachromosomal expression of the antibody encoding nucleic acids (Koller and Smithies, Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); Zijlstra et al., Nature 342:435-438 (1989). In specific embodiments, the expressed antibody molecule is a single chain antibody; alternatively, the nucleic acid sequences include sequences encoding both the heavy and light chains, or fragments thereof, of the antibody.

[0310] Delivery of the nucleic acids into a patient may be either direct, in which case the patient is directly exposed to the nucleic acid or nucleic acid- carrying vectors, or indirect, in which case, cells are first transformed with the nucleic acids in vitro, then transplanted into the patient. These two approaches are known, respectively, as in vivo or ex vivo gene therapy.

[0311] In a specific embodiment, the nucleic acid sequences are directly administered in vivo, where it is expressed to produce the encoded product. This can

be accomplished by any of numerous methods known in the art, e.g., by constructing them as part of an appropriate nucleic acid expression vector and administering it so that they become intracellular, e.g., by infection using defective or attenuated retrovirals or other viral vectors (see U.S. Patent No. 4,980,286), or by direct injection of naked DNA, or by use of microparticle bombardment (e.g., a gene gun; Biolistic, Dupont), or coating with lipids or cell-surface receptors or transfecting agents, encapsulation in liposomes, microparticles, or microcapsules, or by administering them in linkage to a peptide which is known to enter the nucleus, by administering it in linkage to a ligand subject to receptor-mediated endocytosis (see, e.g., Wu and Wu, J. Biol. Chem. 262:4429-4432 (1987)) (which can be used to target cell types specifically expressing the receptors), etc. In another embodiment, nucleic acidligand complexes can be formed in which the ligand comprises a fusogenic viral peptide to disrupt endosomes, allowing the nucleic acid to avoid degradation. In yet another embodiment, the nucleic acid can be targeted in vivo for cell specific uptake and expression, by targeting a specific receptor (see, e.g., PCT Publications WO 92/06180; WO 92/22635; WO92/20316; WO93/14188, WO 93/20221). Alternatively, the nucleic acid can be introduced intracellularly and incorporated within host cell DNA for expression, by homologous recombination (Koller and Smithies, Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); Zijlstra et al., Nature 342:435-438 (1989)).

encoding an antibody of the invention are used. For example, a retroviral vector can be used (see Miller et al., Meth. Enzymol. 217:581-599 (1993)). These retroviral vectors contain the components necessary for the correct packaging of the viral genome and integration into the host cell DNA. The nucleic acid sequences encoding the antibody to be used in gene therapy are cloned into one or more vectors, which facilitates delivery of the gene into a patient. More detail about retroviral vectors can be found in Boesen et al., Biotherapy 6:291-302 (1994), which describes the use of a retroviral vector to deliver the mdr1 gene to hematopoietic stem cells in order to make the stem cells more resistant to chemotherapy. Other references illustrating the use of retroviral vectors in gene therapy are: Clowes et al., J. Clin. Invest. 93:644-651 (1994); Kiem et al., Blood 83:1467-1473 (1994); Salmons and Gunzberg, Human

Gene Therapy 4:129-141 (1993); and Grossman and Wilson, Curr. Opin. in Genetics and Devel. 3:110-114 (1993).

Adenoviruses are other viral vectors that can be used in gene therapy. Adenoviruses are especially attractive vehicles for delivering genes to respiratory epithelia. Adenoviruses naturally infect respiratory epithelia where they cause a mild disease. Other targets for adenovirus-based delivery systems are liver, the central nervous system, endothelial cells, and muscle. Adenoviruses have the advantage of being capable of infecting non-dividing cells. Kozarsky and Wilson, Current Opinion in Genetics and Development 3:499-503 (1993) present a review of adenovirus-based gene therapy. Bout et al., Human Gene Therapy 5:3-10 (1994) demonstrated the use of adenovirus vectors to transfer genes to the respiratory epithelia of rhesus monkeys. Other instances of the use of adenoviruses in gene therapy can be found in Rosenfeld et al., Science 252:431-434 (1991); Rosenfeld et al., Cell 68:143- 155 (1992); Mastrangeli et al., J. Clin. Invest. 91:225-234 (1993); PCT Publication WO94/12649; and Wang, et al., Gene Therapy 2:775-783 (1995). In a preferred embodiment, adenovirus vectors are used.

[0314] Adeno-associated virus (AAV) has also been proposed for use in gene therapy (Walsh et al., Proc. Soc. Exp. Biol. Med. 204:289-300 (1993); U.S. Patent No. 5,436,146).

[0315] Another approach to gene therapy involves transferring a gene to cells in tissue culture by such methods as electroporation, lipofection, calcium phosphate mediated transfection, or viral infection. Usually, the method of transfer includes the transfer of a selectable marker to the cells. The cells are then placed under selection to isolate those cells that have taken up and are expressing the transferred gene. Those cells are then delivered to a patient.

[0316] In this embodiment, the nucleic acid is introduced into a cell prior to administration in vivo of the resulting recombinant cell. Such introduction can be carried out by any method known in the art, including but not limited to transfection, electroporation, microinjection, infection with a viral or bacteriophage vector containing the nucleic acid sequences, cell fusion, chromosome-mediated gene transfer, microcell-mediated gene transfer, spheroplast fusion, etc. Numerous techniques are known in the art for the introduction of foreign genes into cells (see,

e.g., Loeffler and Behr, Meth. Enzymol. 217:599-618 (1993); Cohen et al., Meth. Enzymol. 217:618-644 (1993); Cline, Pharmac. Ther. 29:69-92m (1985) and may be used in accordance with the present invention, provided that the necessary developmental and physiological functions of the recipient cells are not disrupted. The technique should provide for the stable transfer of the nucleic acid to the cell, so that the nucleic acid is expressible by the cell and preferably heritable and expressible by its cell progeny.

- [0317] The resulting recombinant cells can be delivered to a patient by various methods known in the art. Recombinant blood cells (e.g., hematopoietic stem or progenitor cells) are preferably administered intravenously. The amount of cells envisioned for use depends on the desired effect, patient state, etc., and can be determined by one skilled in the art.
- [0318] Cells into which a nucleic acid can be introduced for purposes of gene therapy encompass any desired, available cell type, and include but are not limited to epithelial cells, endothelial cells, keratinocytes, fibroblasts, muscle cells, hepatocytes; blood cells such as T lymphocytes, B lymphocytes, monocytes, macrophages, neutrophils, eosinophils, megakaryocytes, granulocytes; various stem or progenitor cells, in particular hematopoietic stem or progenitor cells, e.g., as obtained from bone marrow, umbilical cord blood, peripheral blood, fetal liver, etc.
- [0319] In a preferred embodiment, the cell used for gene therapy is autologous to the patient.
- [0320] In an embodiment in which recombinant cells are used in gene therapy, nucleic acid sequences encoding an antibody are introduced into the cells such that they are expressible by the cells or their progeny, and the recombinant cells are then administered in vivo for therapeutic effect. In a specific embodiment, stem or progenitor cells are used. Any stem and/or progenitor cells which can be isolated and maintained in vitro can potentially be used in accordance with this embodiment of the present invention (see e.g. PCT Publication WO 94/08598; Stemple and Anderson, Cell 71:973-985 (1992); Rheinwald, Meth. Cell Bio. 21A:229 (1980); and Pittelkow and Scott, Mayo Clinic Proc. 61:771 (1986)).
- [0321] In a specific embodiment, the nucleic acid to be introduced for purposes of gene therapy comprises an inducible promoter operably linked to the coding region,

such that expression of the nucleic acid is controllable by the presence or absence of an appropriate inducer of transcription.

Demonstration of Therapeutic or Prophylactic Activity

preferably tested *in vitro*, and then *in vivo* for the desired therapeutic or prophylactic activity, prior to use in humans. For example, *in vitro* assays to demonstrate the therapeutic or prophylactic utility of a compound or pharmaceutical composition include, the effect of a compound on a cell line or a patient tissue sample. The effect of the compound or composition on the cell line and/or tissue sample can be determined utilizing techniques known to those of skill in the art including, but not limited to, rosette formation assays and cell lysis assays. In accordance with the invention, *in vitro* assays which can be used to determine whether administration of a specific compound is indicated, include *in vitro* cell culture assays in which a patient tissue sample is grown in culture, and exposed to or otherwise administered a compound, and the effect of such compound upon the tissue sample is observed.

Therapeutic/Prophylactic Administration and Composition

[0323] The invention provides methods of treatment, inhibition and prophylaxis by administration to a subject of an effective amount of a compound or pharmaceutical composition of the invention, preferably a polypeptide or antibody of the invention. In a preferred embodiment, the compound is substantially purified (e.g., substantially free from substances that limit its effect or produce undesired side-effects). The subject is preferably an animal, including but not limited to animals such as cows, pigs, horses, chickens, cats, dogs, etc., and is preferably a mammal, and most preferably human.

[0324] Formulations and methods of administration that can be employed when the compound comprises a nucleic acid or an immunoglobulin are described above; additional appropriate formulations and routes of administration can be selected from among those described herein below.

[0325] Various delivery systems are known and can be used to administer a compound of the invention, e.g., encapsulation in liposomes, microparticles.

microcapsules, recombinant cells capable of expressing the compound, receptor-mediated endocytosis (see, e.g., Wu and Wu, J. Biol. Chem. 262:4429-4432 (1987)), construction of a nucleic acid as part of a retroviral or other vector, etc. Methods of introduction include but are not limited to intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, intranasal, epidural, and oral routes. The compounds or compositions may be administered by any convenient route, for example by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (e.g., oral mucosa, rectal and intestinal mucosa, etc.) and may be administered together with other biologically active agents. Administration can be systemic or local. In addition, it may be desirable to introduce the pharmaceutical compounds or compositions of the invention into the central nervous system by any suitable route, including intraventricular and intrathecal injection; intraventricular injection may be facilitated by an intraventricular catheter, for example, attached to a reservoir, such as an Ommaya reservoir. Pulmonary administration can also be employed, e.g., by use of an inhaler or nebulizer, and formulation with an aerosolizing agent.

[0326] In a specific embodiment, it may be desirable to administer the pharmaceutical compounds or compositions of the invention locally to the area in need of treatment; this may be achieved by, for example, and not by way of limitation, local infusion during surgery, topical application, e.g., in conjunction with a wound dressing after surgery, by injection, by means of a catheter, by means of a suppository, or by means of an implant, said implant being of a porous, non-porous, or gelatinous material, including membranes, such as sialastic membranes, or fibers. Preferably, when administering a protein, including an antibody, of the invention, care must be taken to use materials to which the protein does not absorb.

[0327] In another embodiment, the compound or composition can be delivered in a vesicle, in particular a liposome (see Langer, Science 249:1527-1533 (1990); Treat et al., in Liposomes in the Therapy of Infectious Disease and Cancer, Lopez-Berestein and Fidler (eds.), Liss, New York, pp. 353- 365 (1989); Lopez-Berestein, ibid., pp. 317-327; see generally ibid.)

[0328] In yet another embodiment, the compound or composition can be delivered in a controlled release system. In one embodiment, a pump may be used (see Langer, supra; Sefton, CRC Crit. Ref. Biomed. Eng. 14:201 (1987); Buchwald et al., Surgery

88:507 (1980); Saudek et al., N. Engl. J. Med. 321:574 (1989)). In another embodiment, polymeric materials can be used (see Medical Applications of Controlled Release, Langer and Wise (eds.), CRC Pres., Boca Raton, Florida (1974); Controlled Drug Bioavailability, Drug Product Design and Performance, Smolen and Ball (eds.), Wiley, New York (1984); Ranger and Peppas, J., Macromol. Sci. Rev. Macromol. Chem. 23:61 (1983); see also Levy et al., Science 228:190 (1985); During et al., Ann. Neurol. 25:351 (1989); Howard et al., J.Neurosurg. 71:105 (1989)). In yet another embodiment, a controlled release system can be placed in proximity of the therapeutic target, e.g., the brain, thus requiring only a fraction of the systemic dose (see, e.g., Goodson, in Medical Applications of Controlled Release, *supra*, vol. 2, pp. 115-138 (1984)).

[0329] Other controlled release systems are discussed in the review by Langer (Science 249:1527-1533 (1990)).

[0330] In a specific embodiment where the compound of the invention is a nucleic acid encoding a protein, the nucleic acid can be administered *in vivo* to promote expression of its encoded protein, by constructing it as part of an appropriate nucleic acid expression vector and administering it so that it becomes intracellular, e.g., by use of a retroviral vector (see U.S. Patent No. 4,980,286), or by direct injection, or by use of microparticle bombardment (e.g., a gene gun; Biolistic, Dupont), or coating with lipids or cell-surface receptors or transfecting agents, or by administering it in linkage to a homeobox-like peptide which is known to enter the nucleus (see e.g., Joliot et al., Proc. Natl. Acad. Sci. USA 88:1864-1868 (1991)), etc. Alternatively, a nucleic acid can be introduced intracellularly and incorporated within host cell DNA for expression, by homologous recombination.

[0331] The present invention also provides pharmaceutical compositions. Such compositions comprise a therapeutically effective amount of a compound, and a pharmaceutically acceptable carrier. In a specific embodiment, the term "pharmaceutically acceptable" means approved by a regulatory agency of the Federal or a state government or listed in the U.S. Pharmacopeia or other generally recognized pharmacopeia for use in animals, and more particularly in humans. The term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the therapeutic is administered. Such pharmaceutical carriers can be sterile liquids, such as water and

oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil and the like. Water is a preferred carrier when the pharmaceutical composition is administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as liquid carriers, particularly for injectable solutions. Suitable pharmaceutical excipients include starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene, glycol, water, ethanol and the like. The composition, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering agents. These compositions can take the form of solutions, suspensions, emulsion, tablets, pills, capsules, powders, sustained-release formulations and the like. The composition can be formulated as a suppository, with traditional binders and carriers such as triglycerides. Oral formulation can include standard carriers such as pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, etc. Examples of suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences" by E.W. Martin. Such compositions will contain a therapeutically effective amount of the compound, preferably in purified form, together with a suitable amount of carrier so as to provide the form for proper administration to the patient. The formulation should suit the mode of administration.

[0332] In a preferred embodiment, the composition is formulated in accordance with routine procedures as a pharmaceutical composition adapted for intravenous administration to human beings. Typically, compositions for intravenous administration are solutions in sterile isotonic aqueous buffer. Where necessary, the composition may also include a solubilizing agent and a local anesthetic such as lignocaine to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a dry lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent. Where the composition is to be administered by infusion, it can be dispensed with an infusion bottle containing sterile pharmaceutical grade water or saline. Where the composition

is administered by injection, an ampoule of sterile water for injection or saline can be provided so that the ingredients may be mixed prior to administration.

[0333] The compounds of the invention can be formulated as neutral or salt forms. Pharmaceutically acceptable salts include those formed with anions such as those derived from hydrochloric, phosphoric, acetic, oxalic, tartaric acids, etc., and those formed with cations such as those derived from sodium, potassium, ammonium, calcium, ferric hydroxides, isopropylamine, triethylamine, 2-ethylamino ethanol, histidine, procaine, etc.

The amount of the compound of the invention which will be effective in the treatment, inhibition and prevention of a disease or disorder associated with aberrant expression and/or activity of a polypeptide of the invention can be determined by standard clinical techniques. In addition, in vitro assays may optionally be employed to help identify optimal dosage ranges. The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the disease or disorder, and should be decided according to the judgment of the practitioner and each patient's circumstances. Effective doses may be extrapolated from dose-response curves derived from in vitro or animal model test systems.

[0335] For antibodies, the dosage administered to a patient is typically 0.1 mg/kg to 100 mg/kg of the patient's body weight. Preferably, the dosage administered to a patient is between 0.1 mg/kg and 20 mg/kg of the patient's body weight, more preferably 1 mg/kg to 10 mg/kg of the patient's body weight. Generally, human antibodies have a longer half-life within the human body than antibodies from other species due to the immune response to the foreign polypeptides. Thus, lower dosages of human antibodies and less frequent administration is often possible. Further, the dosage and frequency of administration of antibodies of the invention may be reduced by enhancing uptake and tissue penetration (e.g., into the brain) of the antibodies by modifications such as, for example, lipidation.

[0336] The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Optionally associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture,

use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

Diagnosis and Imaging

Labeled antibodies, and derivatives and analogs thereof, which specifically bind to a polypeptide of interest can be used for diagnostic purposes to detect, diagnose, or monitor diseases, disorders, and/or conditions associated with the aberrant expression and/or activity of a polypeptide of the invention. The invention provides for the detection of aberrant expression of a polypeptide of interest, comprising (a) assaying the expression of the polypeptide of interest in cells or body fluid of an individual using one or more antibodies specific to the polypeptide interest and (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of aberrant expression.

The invention provides a diagnostic assay for diagnosing a reproductive system disorder, comprising (a) assaying the expression of the polypeptide of interest in cells or body fluid of an individual using one or more antibodies specific to the polypeptide interest and (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a particular disorder. With respect to cancer, the presence of a relatively high amount of transcript in biopsied tissue from an individual may indicate a predisposition for the development of the disease, or may provide a means for detecting the disease prior to the appearance of actual clinical symptoms. A more definitive diagnosis of this type may allow health professionals to employ preventative measures or aggressive treatment earlier thereby preventing the development or further progression of the cancer.

[0339] Antibodies of the invention can be used to assay protein levels in a biological sample using classical immunohistological methods known to those of skill in the art (e.g., see Jalkanen et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen et al., J. Cell . Biol. 105:3087-3096 (1987)). Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked

immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase; radioisotopes, such as iodine (125I, 121I), carbon (14C), sulfur (35S), tritium (3H), indium (112In), and technetium (99Tc); luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin.

One facet of the invention is the detection and diagnosis of a disease or [0340] disorder associated with aberrant expression of a polypeptide of interest in an animal, preferably a mammal and most preferably a human. A preferred embodiment of the invention is the detection and diagnosis of a disease or disorder of the reproductive system associated with aberrant expression of a reproductive system antigen in an animal, preferably a mammal and most preferably a human. In one embodiment, diagnosis comprises: a) administering (for example, parenterally, subcutaneously, or intraperitoneally) to a subject an effective amount of a labeled molecule which specifically binds to the polypeptide of interest; b) waiting for a time interval following the administering for permitting the labeled molecule to preferentially concentrate at sites in the subject where the polypeptide is expressed (and for unbound labeled molecule to be cleared to background level); c) determining background level; and d) detecting the labeled molecule in the subject, such that detection of labeled molecule above the background level indicates that the subject has a particular disease or disorder associated with aberrant expression of the polypeptide of interest. Background level can be determined by various methods including, comparing the amount of labeled molecule detected to a standard value previously determined for a particular system.

[0341] It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of 99mTc. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the specific protein. *In vivo* tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in Tumor Imaging: The

Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)).

[0342] Depending on several variables, including the type of label used and the mode of administration, the time interval following the administration for permitting the labeled molecule to preferentially concentrate at sites in the subject and for unbound labeled molecule to be cleared to background level is 6 to 48 hours or 6 to 24 hours or 6 to 12 hours. In another embodiment the time interval following administration is 5 to 20 days or 5 to 10 days.

[0343] In an embodiment, monitoring of the disease or disorder is carried out by repeating the method for diagnosing the disease or disorder, for example, one month after initial diagnosis, six months after initial diagnosis, one year after initial diagnosis, etc.

[0344] Presence of the labeled molecule can be detected in the patient using methods known in the art for *in vivo* scanning. These methods depend upon the type of label used. Skilled artisans will be able to determine the appropriate method for detecting a particular label. Methods and devices that may be used in the diagnostic methods of the invention include, but are not limited to, computed tomography (CT), whole body scan such as position emission tomography (PET), magnetic resonance imaging (MRI), and sonography.

In a specific embodiment, the molecule is labeled with a radioisotope and is detected in the patient using a radiation responsive surgical instrument (Thurston et al., U.S. Patent No. 5,441,050). In another embodiment, the molecule is labeled with a fluorescent compound and is detected in the patient using a fluorescence responsive scanning instrument. In another embodiment, the molecule is labeled with a positron emitting metal and is detected in the patent using positron emission-tomography. In yet another embodiment, the molecule is labeled with a paramagnetic label and is detected in a patient using magnetic resonance imaging (MRI).

Kits

[0346] The present invention provides kits that can be used in the above methods. In one embodiment, a kit comprises an antibody of the invention, preferably a purified antibody, in one or more containers. In a specific embodiment, the kits of the present

invention contain a substantially isolated polypeptide comprising an epitope which is specifically immunoreactive with an antibody included in the kit. Preferably, the kits of the present invention further comprise a control antibody which does not react with the polypeptide of interest. In another specific embodiment, the kits of the present invention contain a means for detecting the binding of an antibody to a polypeptide of interest (e.g., the antibody may be conjugated to a detectable substrate such as a fluorescent compound, an enzymatic substrate, a radioactive compound or a luminescent compound, or a second antibody which recognizes the first antibody may be conjugated to a detectable substrate).

In another specific embodiment of the present invention, the kit is a diagnostic kit for use in screening serum containing antibodies specific against proliferative and/or cancerous polynucleotides and polypeptides. Such a kit may include a control antibody that does not react with the polypeptide of interest. Such a kit may include a substantially isolated polypeptide antigen comprising an epitope which is specifically immunoreactive with at least one anti-polypeptide antigen antibody. Further, such a kit includes means for detecting the binding of said antibody to the antigen (e.g., the antibody may be conjugated to a fluorescent compound such as fluorescein or rhodamine which can be detected by flow cytometry). In specific embodiments, the kit may include a recombinantly produced or chemically synthesized polypeptide antigen. The polypeptide antigen of the kit may also be attached to a solid support.

[0348] In a more specific embodiment the detecting means of the above-described kit includes a solid support to which said polypeptide antigen is attached. Such a kit may also include a non-attached reporter-labeled anti-human antibody. In this embodiment, binding of the antibody to the polypeptide antigen can be detected by binding of the said reporter-labeled antibody.

[0349] In an additional embodiment, the invention includes a diagnostic kit for use in screening serum containing antigens of the polypeptide of the invention. The diagnostic kit includes a substantially isolated antibody specifically immunoreactive with polypeptide or polynucleotide antigens, and means for detecting the binding of the polynucleotide or polypeptide antigen to the antibody. In one embodiment, the antibody is attached to a solid support. In a specific embodiment, the antibody may be

a monoclonal antibody. The detecting means of the kit may include a second, labeled monoclonal antibody. Alternatively, or in addition, the detecting means may include a labeled, competing antigen.

reagent having a surface-bound antigen obtained by the methods of the present invention. After binding with specific antigen antibody to the reagent and removing unbound serum components by washing, the reagent is reacted with reporter-labeled anti-human antibody to bind reporter to the reagent in proportion to the amount of bound anti-antigen antibody on the solid support. The reagent is again washed to remove unbound labeled antibody, and the amount of reporter associated with the reagent is determined. Typically, the reporter is an enzyme which is detected by incubating the solid phase in the presence of a suitable fluorometric, luminescent or colorimetric substrate (Sigma, St. Louis, MO).

[0351] The solid surface reagent in the above assay is prepared by known techniques for attaching protein material to solid support material, such as polymeric beads, dip sticks, 96-well plate or filter material. These attachment methods generally include non-specific adsorption of the protein to the support or covalent attachment of the protein, typically through a free amine group, to a chemically reactive group on the solid support, such as an activated carboxyl, hydroxyl, or aldehyde group. Alternatively, streptavidin coated plates can be used in conjunction with biotinylated antigen(s).

[0352] Thus, the invention provides an assay system or kit for carrying out this diagnostic method. The kit generally includes a support with surface-bound recombinant antigens, and a reporter-labeled anti-human antibody for detecting surface-bound anti-antigen antibody.

Uses of the Polynucleotides

[0353] Each of the polynucleotides identified herein can be used in numerous ways as reagents. The following description should be considered exemplary and utilizes known techniques.

The polynucleotides of the present invention are useful for chromosome identification. There exists an ongoing need to identify new chromosome markers, since few chromosome marking reagents, based on actual sequence data (repeat polymorphisms), are presently available. Each sequence is specifically targeted to and can hybridize with a particular location on an individual human chromosome, thus each polynucleotide of the present invention can routinely be used as a chromosome marker using techniques known in the art. Table 1A, column 8 provides the chromosome location of some of the polynucleotides of the invention.

primers (preferably at least 15 bp (e.g., 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can optionally be selected using computer analysis so that primers do not span more than one predicted exon in the genomic DNA. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to SEQ ID NO:X will yield an amplified fragment.

[0356] Similarly, somatic hybrids provide a rapid method of PCR mapping the polynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can be achieved with panels of specific chromosome fragments. Other gene mapping strategies that can be used include in situ hybridization, prescreening with labeled flow-sorted chromosomes, preselection by hybridization to construct chromosome specific-cDNA libraries, and computer mapping techniques (See, e.g., Shuler, Trends Biotechnol 16:456-459 (1998) which is hereby incorporated by reference in its entirety).

[0357] Precise chromosomal location of the polynucleotides can also be achieved using fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides 2,000-4,000 bp are preferred. For a review of this technique, see Verma et al., "Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York (1988).

[0358] For chromosome mapping, the polynucleotides can be used individually (to mark a single chromosome or a single site on that chromosome) or in panels (for marking multiple sites and/or multiple chromosomes).

[0359] Thus, the present invention also provides a method for chromosomal localization which involves (a) preparing PCR primers from the polynucleotide sequences in Table 1A and/or Table 2 and SEQ ID NO:X and (b) screening somatic cell hybrids containing individual chromosomes.

[0360] The polynucleotides of the present invention would likewise be useful for radiation hybrid mapping, HAPPY mapping, and long range restriction mapping. For a review of these techniques and others known in the art, see, e.g. Dear, "Genome Mapping: A Practical Approach," IRL Press at Oxford University Press, London (1997); Aydin, J. Mol. Med. 77:691-694 (1999); Hacia et al., Mol. Psychiatry 3:483-492 (1998); Herrick et al., Chromosome Res. 7:409-423 (1999); Hamilton et al., Methods Cell Biol. 62:265-280 (2000); and/or Ott, J. Hered. 90:68-70 (1999), each of which is hereby incorporated by reference in its entirety.

[0361] Once a polynucleotide has been mapped to a precise chromosomal location, the physical position of the polynucleotide can be used in linkage analysis. Linkage analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. (Disease mapping data are found, for example, in V. McKusick, Mendelian Inheritance in Man (available on line through Johns Hopkins University Welch Medical Library).) Column 9 of Table 1A provides an OMIM reference identification number of diseases associated with the cytologic band disclosed in column 8 of Table 1A, as determined using techniques described herein and by reference to Table 5. Assuming 1 megabase mapping resolution and one gene per 20 kb, a cDNA precisely localized to a chromosomal region associated with the disease could be one of 50-500 potential causative genes.

[0362] Thus, once coinheritance is established, differences in a polynucleotide of the invention and the corresponding gene between affected and unaffected individuals can be examined. First, visible structural alterations in the chromosomes, such as deletions or translocations, are examined in chromosome spreads or by PCR. If no structural alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected individuals, but not in normal individuals, indicates

that mutations may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal individuals is required to distinguish the mutation from a polymorphism. If a new polymorphism is identified, this polymorphic polypeptide can be used for further linkage analysis.

[0363] Furthermore, increased or decreased expression of the gene in affected individuals as compared to unaffected individuals can be assessed using the polynucleotides of the invention. Any of these alterations (altered expression, chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic marker. Diagnostic and prognostic methods, kits and reagents encompassed by the present invention are briefly described below and more thoroughly elsewhere herein (see e.g., the sections labeled "Antibodies", "Diagnostic Assays", and "Methods for Detecting Reproductive System Disease, Including Cancer").

[0364] Thus, the invention also provides a diagnostic method useful during diagnosis of a disorder, involving measuring the expression level of polynucleotides of the present invention in cells or body fluid from an individual and comparing the measured gene expression level with a standard level of polynucleotide expression level, whereby an increase or decrease in the gene expression level compared to the standard is indicative of a disorder. Additional non-limiting examples of diagnostic methods encompassed by the present invention are more thoroughly described elsewhere herein (see, e.g., Example 12).

[0365] In still another embodiment, the invention includes a kit for analyzing samples for the presence of proliferative and/or cancerous polynucleotides derived from a test subject, as further described herein. In a general embodiment, the kit includes at least one polynucleotide probe containing a nucleotide sequence that will specifically hybridize with a polynucleotide of the invention and a suitable container. In a specific embodiment, the kit includes two polynucleotide probes defining an internal region of the polynucleotide of the invention, where each probe has one strand containing a 31'mer-end internal to the region. In a further embodiment, the probes may be useful as primers for polymerase chain reaction amplification.

[0366] Where a diagnosis of a related disorder, including, for example, diagnosis of a tumor, has already been made according to conventional methods, the present invention is useful as a prognostic indicator, whereby patients exhibiting enhanced or

depressed polynucleotide of the invention expression will experience a worse clinical outcome relative to patients expressing the gene at a level nearer the standard level.

By "measuring the expression level of polynucleotides of the invention" is intended qualitatively or quantitatively measuring or estimating the level of the polypeptide of the invention or the level of the mRNA encoding the polypeptide of the invention in a first biological sample either directly (e.g., by determining or estimating absolute protein level or mRNA level) or relatively (e.g., by comparing to the polypeptide level or mRNA level in a second biological sample). Preferably, the polypeptide level or mRNA level in the first biological sample is measured or estimated and compared to a standard polypeptide level or mRNA level, the standard being taken from a second biological sample obtained from an individual not having the related disorder or being determined by averaging levels from a population of individuals not having a related disorder. As will be appreciated in the art, once a standard polypeptide level or mRNA level is known, it can be used repeatedly as a standard for comparison.

[0368] By "biological sample" is intended any biological sample obtained from an individual, body fluid, cell line, tissue culture, or other source which contains polypeptide of the present invention or the corresponding mRNA. As indicated, biological samples include body fluids (such as semen, lymph, vaginal pool, sera, plasma, urine, synovial fluid and spinal fluid) which contain the polypeptide of the present invention, and tissue sources found to express the polypeptide of the present invention. Methods for obtaining tissue biopsies and body fluids from mammals are well known in the art. Where the biological sample is to include mRNA, a tissue biopsy is the preferred source.

[0369] The method(s) provided above may preferably be applied in a diagnostic method and/or kits in which polynucleotides and/or polypeptides of the invention are attached to a solid support. In one exemplary method, the support may be a "gene chip" or a "biological chip" as described in U.S. Patents 5,837,832, 5,874,219, and 5,856,174. Further, such a gene chip with polynucleotides of the invention attached may be used to identify polymorphisms between the isolated polynucleotide sequences of the invention, with polynucleotides isolated from a test subject. The knowledge of such polymorphisms (i.e., their location, as well as, their existence) would be

beneficial in identifying disease loci for many disorders, such as for example, in neural disorders, immune system disorders, muscular disorders, reproductive disorders, gastrointestinal disorders, pulmonary disorders, digestive disorders, cardiovascular disorders, renal disorders, proliferative disorders, and/or cancerous diseases and conditions. Such a method is described in U.S. Patents 5,858,659 and 5,856,104. The U.S. Patents referenced *supra* are hereby incorporated by reference in their entirety herein.

[0370] The present invention encompasses polynucleotides of the present invention that are chemically synthesized, or reproduced as peptide nucleic acids (PNA), or according to other methods known in the art. The use of PNAs would serve as the preferred form if the polynucleotides of the invention are incorporated onto a solid support, or gene chip. For the purposes of the present invention, a peptide nucleic acid (PNA) is a polyamide type of DNA analog and the monomeric units for adenine, guanine, thymine and cytosine are available commercially (Perceptive Biosystems). Certain components of DNA, such as phosphorus, phosphorus oxides, or deoxyribose derivatives, are not present in PNAs. As disclosed by Nielsen et al., Science 254:1497 (1991); and Egholm et al., Nature 365:666 (1993), PNAs bind specifically and tightly to complementary DNA strands and are not degraded by nucleases. In fact, PNA binds more strongly to DNA than DNA itself does. This is probably because there is no electrostatic repulsion between the two strands, and also the polyamide backbone is more flexible. Because of this, PNA/DNA duplexes bind under a wider range of stringency conditions than DNA/DNA duplexes, making it easier to perform multiplex hybridization. Smaller probes can be used than with DNA due to the strong binding. In addition, it is more likely that single base mismatches can be determined with PNA/DNA hybridization because a single mismatch in a PNA/DNA 15-mer lowers the melting point (T.sub.m) by 8°-20° C, vs. 4°-16° C for the DNA/DNA 15-mer duplex. Also, the absence of charge groups in PNA means that hybridization can be done at low ionic strengths and reduce possible interference by salt during the analysis.

[0371] The compounds of the present invention have uses which include, but are not limited to, detecting cancer in mammals. In particular the invention is useful during diagnosis of pathological cell proliferative neoplasias which include, but are not limited to: acute myelogenous leukemias including acute monocytic leukemia,

acute myeloblastic leukemia, acute promyelocytic leukemia, acute myelomonocytic leukemia, acute erythroleukemia, acute megakaryocytic leukemia, and acute undifferentiated leukemia, etc.; and chronic myelogenous leukemias including chronic myelomonocytic leukemia, chronic granulocytic leukemia, etc. Preferred mammals include monkeys, apes, cats, dogs, cows, pigs, horses, rabbits and humans. Particularly preferred are humans.

[0372] The compounds of the present invention have preferred uses which include, but are not limited to, detecting reproductive system cancers in mammals. particular the invention is useful during diagnosis of pathological cell proliferative neoplasias which include, but are not limited to: testicular cancers (including, for example, teratocarcinoma, embryonal cell carcinoma, yolk sac tumors, Leydig cell tumors, and as listed below in the section entitiled "Reproductive System Disorders"), prostate cancers (e.g., adenocarcinomas, transitional cell carcinomas, ductal carcinomas, squamous cell carcinomas, and as listed below in the section entitled "Reproductive System Disorders"), penile cancers (such as squamous cell carcinoma, verrucous carcinoma, penile urethral carcinoma, and as listed below in the section entitled "Reproductive System Disorders"), cancers of the vagina and vulva (including, for example, basal cell carcinoma, melanomas, cancer of Bartholin's gland, and as listed below in the section entitled "Reproductive System Disorders"), uterine cancers (such as adenocarcinomas, keiomyosarcomas, and as listed below in the section entitled "Reproductive System Disorders"), ovarian cancers (e.g., Sertoli-Leydig tumors, endometriod carcinoma of the ovary, ovarian papollary serous adenocarcinoma, ovarian Krukenberg tumors, and as listed below in the section entitled "Reproductive System Disorders"), and cancers of the cervix (including, for example, squamous metaplasia, columnar cell neoplasia, squamous cell carcinoma, and as listed below in the section entitled "Reproductive System Disorders"). Preferred mammals include monkeys, apes, cats, dogs, cows, pigs, horses, rabbits and humans. Particularly preferred are humans.

[0373] Pathological cell proliferative disorders are often associated with inappropriate activation of proto-oncogenes. (Gelmann, E. P. et al., "The Etiology of Acute Leukemia: Molecular Genetics and Viral Oncology," in Neoplastic Diseases of the Blood, Vol 1., Wiernik, P. H. et al. eds., 161-182 (1985)). Neoplasias are now

believed to result from the qualitative alteration of a normal cellular gene product, or from the quantitative modification of gene expression by insertion into the chromosome of a viral sequence, by chromosomal translocation of a gene to a more actively transcribed region, or by some other mechanism. (Gelmann et al., supra) It is likely that mutated or altered expression of specific genes is involved in the pathogenesis of some leukemias, among other tissues and cell types. (Gelmann et al., supra) Indeed, the human counterparts of the oncogenes involved in some animal neoplasias have been amplified or translocated in some cases of human leukemia and carcinoma. (Gelmann et al., supra)

[0374] For example, c-myc expression is highly amplified in the non-lymphocytic leukemia cell line HL-60. When HL-60 cells are chemically induced to stop proliferation, the level of c-myc is found to be downregulated. (International Publication Number WO 91/15580). However, it has been shown that exposure of HL-60 cells to a DNA construct that is complementary to the 5' end of c-myc or c-myb blocks translation of the corresponding mRNAs which downregulates expression of the c-myc or c-myb proteins and causes arrest of cell proliferation and differentiation of the treated cells. (International Publication Number WO 91/15580; Wickstrom et al., Proc. Natl. Acad. Sci. 85:1028 (1988); Anfossi et al., Proc. Natl. Acad. Sci. 86:3379 (1989)). However, the skilled artisan would appreciate the present invention's usefulness is not be limited to treatment, prevention, diagnosis and/or prognosis, of proliferative disorders of cells and tissues of hematopoietic origin, in light of the numerous cells and cell types of varying origins which are known to exhibit proliferative phenotypes. In preferred embodiments, the compounds and/or methods of the invention are used to treat, prevent, diagnose, and/or prognose, proliferative disorders of reproductive system cells and tissues.

[0375] In addition to the foregoing, a polynucleotide of the present invention can be used to control gene expression through triple helix formation or through antisense DNA or RNA. Antisense techniques are discussed, for example, in Okano, J. Neurochem. 56: 560 (1991); "Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988). Triple helix formation is discussed in, for instance Lee et al., Nucleic Acids Research 6: 3073 (1979); Cooney et al., Science 241: 456 (1988); and Dervan et al., Science 251: 1360 (1991). Both methods rely on

binding of the polynucleotide to a complementary DNA or RNA. techniques, preferred polynucleotides are usually oligonucleotides 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxy-nucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988).) Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. The oligonucleotide described above can also be delivered to cells such that the antisense RNA or DNA may be expressed in vivo to inhibit production of polypeptide of the present invention antigens. Both techniques are effective in model systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat disease, and in particular, for the treatment of proliferative diseases and/or conditions. Non-limiting antisense and triple helix methods encompassed by the present invention are more thoroughly described elsewhere herein (see, e.g., the section labeled "Antisense and Ribozyme (Antagonists)").

One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the present invention offer a means of targeting such genetic defects in a highly accurate manner. Another goal is to insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell. Additional non-limiting examples of gene therapy methods encompassed by the present invention are more thoroughly described elsewhere herein (see, e.g., the sections labeled "Gene Therapy Methods" and Examples 16, 17 and 18).

[0377] The polynucleotides are also useful for identifying individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for

identifying personnel. This method does not suffer from the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The polynucleotides of the present invention can be used as additional DNA markers for RFLP.

The polynucleotides of the present invention can also be used as an alternative to RFLP, by determining the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, individuals can be identified because each individual will have a unique set of DNA sequences. Once an unique ID database is established for an individual, positive identification of that individual, living or dead, can be made from extremely small tissue samples.

[0379] Forensic biology also benefits from using DNA-based identification techniques as disclosed herein. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, synovial fluid, amniotic fluid, breast milk, lymph, pulmonary sputum or surfactant, urine, fecal matter, etc., can be amplified using PCR. In one prior art technique, gene sequences amplified from polymorphic loci, such as DQa class II HLA gene, are used in forensic biology to identify individuals. (Erlich, H., PCR Technology, Freeman and Co. (1992).) Once these specific polymorphic loci are amplified, they are digested with one or more restriction enzymes, yielding an identifying set of bands on a Southern blot probed with DNA corresponding to the DQa class II HLA gene. Similarly, polynucleotides of the present invention can be used as polymorphic markers for forensic purposes.

[0380] There is also a need for reagents capable of identifying the source of a particular tissue. Such need arises, for example, in forensics when presented with tissue of unknown origin. Appropriate reagents can comprise, for example, DNA probes or primers prepared from the sequences of the present invention, specific to tissues, including but not limited to, those sequences referred to in Table 1A. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for contamination. Additional non-limiting examples of such uses are further described herein.

[0381] Because reproductive system antigens are found expressed in reproductive system tissues, the polynucleotides of the present invention are also useful as hybridization probes for differential identification of the tissue(s) or cell type(s) present in a biological sample. Similarly, polypeptides and antibodies directed to polypeptides of the present invention are useful to provide immunological probes for differential identification of the tissue(s) (e.g., immunohistochemistry assays) or cell type(s) (e.g., immunocytochemistry assays). In a specific embodiment, the polynucleotides of the present invention are also useful as hybridization probes for differential identification of reproductive system tissue(s) or cell type(s) present in a biological sample. Similarly, polypeptides and antibodies directed to polypeptides of the present invention are useful to provide immunological probes for differential identification of reproductive system tissue(s) (e.g., immunohistochemistry assays) or cell type(s) (e.g., immunocytochemistry assays). In addition, for a number of disorders of the above tissues or cells, significantly higher or lower levels of gene expression of the polynucleotides/polypeptides of the present invention may be detected in certain tissues (e.g., tissues expressing polypeptides and/or polynucleotides of the present invention, for example, normal reproductive system tissues or diseased reproductive system tissues, and/or those tissues/cells corresponding to the library source relating to a polynucleotide sequence of the invention as disclosed in column 7 of Table 1A, and/or cancerous and/or wounded tissues) or bodily fluids (e.g., semen, lymph, vaginal pool, serum, plasma, urine, synovial fluid or spinal fluid) taken from an individual having such a disorder, relative to a "standard" gene expression level, i.e., the expression level in healthy tissue from an individual not having the disorder.

[0382] Thus, the invention provides a diagnostic method of a disorder, which involves: (a) assaying gene expression level in cells or body fluid of an individual; (b) comparing the gene expression level with a standard gene expression level, whereby an increase or decrease in the assayed gene expression level compared to the standard expression level is indicative of a disorder.

[0383] In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and

making oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response.

Uses of the Polypeptides

[0384] Each of the polypeptides identified herein can be used in numerous ways.

The following description should be considered exemplary and utilizes known techniques.

[0385] Polypeptides and antibodies directed to polypeptides of the present invention are useful to provide immunological probes for differential identification of the tissue(s) (e.g., immunohistochemistry assays such as, for example, ABC immunoperoxidase (Hsu et al., J. Histochem. Cytochem. 29:577-580 (1981)) or cell type(s) (e.g., immunocytochemistry assays).

polynucleotides of the invention in a biological sample using classical immunohistological methods known to those of skill in the art (see, e.g., Jalkanen, et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, et al., J. Cell. Biol. 105:3087-3096 (1987)). Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase; radioisotopes, such as iodine (131 I, 125 I, 123 I, 121 I), carbon (14 C), sulfur (35 S), tritium (3 H), indium (115 m In, 113 m In, 112 In, 111 ln), and technetium (99 Tc, 99 m Tc), thallium (201 Ti), gallium (68 Ga, 67 Ga), palladium (103 Pd), molybdenum (99 Mo), xenon (133 Xe), fluorine (18 F), 153 Sm, 177 Lu, 159 Gd, 149 Pm, 140 La, 175 Yb, 166 Ho, 90 Y, 47 Sc, 186 Re, 188 Re, 142 Pr, 105 Rh, 97 Ru; luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin.

[0387] In addition to assaying levels of polypeptide of the present invention in a biological sample, proteins can also be detected *in vivo* by imaging. Antibody labels or markers for *in vivo* imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable

characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

103881 A reproductive system antigen-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, 131 I, 112 In, 99m Tc, (131 I, 125 I, 123 I, 121 I), carbon (14C), sulfur (35S), tritium (3H), indium (115mIn, 113mIn, 112In, 111In), and technetium (99Tc, 99mTc), thallium (201Ti), gallium (68Ga, 67Ga), palladium (103Pd), molybdenum (99Mo), xenon (133Xe), fluorine (18F, 153Sm, 177Lu, 159Gd, 149Pm, 140La, 175Yb, 166Ho, 90Y, 47Sc, 186Re, ¹⁸⁸Re, ¹⁴²Pr, ¹⁰⁵Rh, ⁹⁷Ru), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously or intraperitoneally) into the mammal to be examined for reproductive system disorder. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of 99mTc. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which express the polypeptide encoded by a polynucleotide of the invention. In vivo tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments" (Chapter 13 in Tumor Imaging: The Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)).

[0389] In one embodiment, the invention provides a method for the specific delivery of compositions of the invention to cells by administering polypeptides of the invention (e.g., polypeptides encoded by polynucleotides of the invention and/or antibodies) that are associated with heterologous polypeptides or nucleic acids. In one example, the invention provides a method for delivering a therapeutic protein into the targeted cell. In another example, the invention provides a method for delivering a single stranded nucleic acid (e.g., antisense or ribozymes) or double stranded nucleic acid (e.g., DNA that can integrate into the cell's genome or replicate episomally and that can be transcribed) into the targeted cell.

[0390] In another embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention in association with toxins or cytotoxic prodrugs.

[0391] In a preferred embodiment, the invention provides a method for the specific destruction of reproductive system cells (e.g., aberrant reproductive system cells, neoplasms of the reproductive system) by administering polypeptides of the invention (e.g., polypeptides encoded by polynucleotides of the invention and/or antibodies) in association with toxins or cytotoxic prodrugs. In another preferred embodiment the invention provides a method for the specific destruction of tissues/cells corresponding to the library source relating to a polynucleotide sequence of the invention as disclosed in column 7 of Table 1A by administering polypeptides of the invention in association with toxins or cytotoxic prodrugs.

[0392] By "toxin" is meant one or more compounds that bind and activate endogenous cytotoxic effector systems, radioisotopes, holotoxins, modified toxins, catalytic subunits of toxins, or any molecules or enzymes not normally present in or on the surface of a cell that under defined conditions cause the cell's death. Toxins that may be used according to the methods of the invention include, but are not limited to, radioisotopes known in the art, compounds such as, for example, antibodies (or complement fixing containing portions thereof) that bind an inherent or induced endogenous cytotoxic effector system, thymidine kinase, endonuclease, RNAse, alpha toxin, ricin, abrin, Pseudomonas exotoxin A, diphtheria toxin, saporin, momordin, gelonin, pokeweed antiviral protein, alpha-sarcin and cholera toxin. "Toxin" also includes a cytostatic or cytocidal agent, a therapeutic agent or a radioactive metal ion, e.g., alpha-emitters such as, for example, ²¹³Bi, or other radioisotopes such as, for example, 103Pd, 133Xe, 131I, 111In, 68Ge, 57Co, 65Zn, 85Sr, 32P, 35S, 90Y, 153Sm, 153Gd, ¹⁶⁹Yb, ⁵¹Cr, ⁵⁴Mn, ⁷⁵Se, ¹¹³Sn, ⁹⁰Yttrium, ¹¹⁷Tin, ¹⁸⁶Rhenium, ¹⁶⁶Holmium, and 188Rhenium; luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin.

[0393] In a specific embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention or antibodies of the invention in association with the radioisotope ⁹⁰Y. In another specific embodiment, the invention provides a method for the specific

destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention or antibodies of the invention in association with the radioisotope ¹¹¹In. In a further specific embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention or antibodies of the invention in association with the radioisotope ¹³¹I.

[0394] Techniques known in the art may be applied to label polypeptides of the invention (including antibodies). Such techniques include, but are not limited to, the use of bifunctional conjugating agents (see e.g., U.S. Patent Nos. 5,756,065; 5,714,631; 5,696,239; 5,652,361; 5,505,931; 5,489,425; 5,435,990; 5,428,139; 5,342,604; 5,274,119; 4,994,560; and 5,808,003; the contents of each of which are hereby incorporated by reference in its entirety).

[0395] Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression level of a polypeptide of the present invention in cells or body fluid of an individual; and (b) comparing the assayed polypeptide expression level with a standard polypeptide expression level, whereby an increase or decrease in the assayed polypeptide expression level compared to the standard expression level is indicative of a disorder. With respect to cancer, the presence of a relatively high amount of transcript in biopsied tissue from an individual may indicate a predisposition for the development of the disease, or may provide a means for detecting the disease prior to the appearance of actual clinical symptoms. A more definitive diagnosis of this type may allow health professionals to employ preventative measures or aggressive treatment earlier thereby preventing the development or further progression of the cancer.

[0396] Moreover, polypeptides of the present invention can be used to treat or prevent diseases or conditions of the reproductive system such as, for example, reproductive system injury and trauma, infections, neoplastic disorders, congenital defects, and diseases or disorders which result in infertility, complications with pregnancy, labor, or parturition, postpartum difficulties, and as listed below in the section entitled "Reproductive System Disorders". In preferred embodiments, polynucleotides expressed in a particular tissue type (see, e.g., Table 1A, column 7) are used to diagnose, detect, prevent, treat and/or prognose disorders associated with

the tissue type. For example, patients can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide (e.g., insulin), to supplement absent or decreased levels of a different polypeptide (e.g., hemoglobin S for hemoglobin B, SOD, catalase, DNA repair proteins), to inhibit the activity of a polypeptide (e.g., an oncogene or tumor supressor), to activate the activity of a polypeptide (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor by competing with it for free ligand (e.g., soluble TNF receptors used in reducing inflammation), or to bring about a desired response (e.g., blood vessel growth inhibition, enhancement of the immune response to proliferative cells or tissues).

[0397] Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat disease (as described *supra*, and elsewhere herein). For example, administration of an antibody directed to a polypeptide of the present invention can bind, and/or neutralize the polypeptide, and/or reduce overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

[0398] At the very least, the polypeptides of the present invention can be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the biological activities described herein.

Diagnostic Asssays

[0399] The compounds of the present invention are useful for diagnosis, treatment, prevention and/or prognosis of various reproductive system related disorders in mammals, preferably humans. Such disorders include, but are not limited to, reproductive system injury and trauma, infections, neoplastic disorders, congenital defects, and diseases or disorders which result in infertility, complications with pregnancy, labor, or parturition, postpartum difficulties, and as listed below in the section entitled "Reproductive System Disorders". In preferred embodiments,

polynucleotides expressed in a particular tissue type (see, e.g., Table 1A, column 7) are used to diagnose, detect, prevent, treat and/or prognose disorders associated with the tissue type.

[0400] Reproductive system antigens are expressed in reproductive system. For a number of reproductive system-related disorders, substantially altered (increased or decreased) levels of reproductive system antigen gene expression can be detected in reproductive system tissue or other cells or bodily fluids (e.g., sera, plasma, urine, semen, synovial fluid or spinal fluid) taken from an individual having such a disorder, relative to a "standard" reproductive system antigen gene expression level, that is, the reproductive system antigen expression level in reproductive system tissues or bodily fluids from an individual not having the reproductive system disorder. Thus, the invention provides a diagnostic method useful during diagnosis of a reproductive system disorder, which involves measuring the expression level of the gene encoding the reproductive system associated polypeptide in reproductive system tissue or other cells or body fluid from an individual and comparing the measured gene expression level with a standard reproductive system antigens gene expression level, whereby an increase or decrease in the gene expression level(s) compared to the standard is indicative of an reproductive system disorder.

during diagnosis of a disorder of a normal or diseased tissue/cell source corresponding to column 7 of Table 1A, which involves measuring the expression level of the coding sequence of a polynucleotide sequence associated with this tissue/cell source as disclosed in Table 1A in the tissue/cell source or other cells or body fluid from an individual and comparing the expression level of the coding sequence with a standard expression level of the coding sequence, whereby an increase or decrease in the gene expression level(s) compared to the standard is indicative of a disorder of a normal or diseased tissue/cell source corresponding to column 7 of Table 1A.

[0402] In particular, it is believed that certain tissues in mammals with cancer of cells or tissue of the reproductive system express significantly enhanced or reduced levels of normal or altered reproductive system antigen expression and mRNA encoding the reproductive system associated polypeptide when compared to a

corresponding "standard" level. Further, it is believed that enhanced or depressed levels of the reproductive system associated polypeptide can be detected in certain body fluids (e.g., sera, plasma, urine, and spinal fluid) or cells or tissue from mammals with such a cancer when compared to sera from mammals of the same species not having the cancer.

For example, as disclosed herein, reproductive system associated [0403] polypeptides of the invention are expressed in tissues of the reproductive system. Accordingly, polynucleotides of the invention (e.g., polynucleotide sequences complementary to all or a portion of a reproductive system antigen mRNA nucleotide sequence of SEQ ID NO:X, nucleotide sequence encoding SEQ ID NO:Y, nucleotide sequence encoding a polypeptide encoded by SEQ ID' NO:X and/or a nucleotide sequence delineated by columns 8 and 9 of Table 2) and antibodies (and antibody fragments) directed against the polypeptides of the invention may be used to quantitate or qualitate concentrations of cells of the reproductive system expressing reproductive system antigens, preferrably on their cell surfaces. These polynucleotides and antibodies additionally have diagnostic applications in detecting abnormalities in the level of reproductive system antigens gene expression, or abnormalities in the structure and/or temporal, tissue, cellular, or subcellular location of reproductive system antigens. These diagnostic assays may be performed in vivo or in vitro, such as, for example, on blood samples, biopsy tissue or autopsy tissue. In specific embodiments, polynucleotides and antibodies of the invention are used to quantitate or qualitate tissues/cells corresponding to the library source disclosed in column 7 of Table 1A expressing the corresponding reproductive system sequence disclosed in the same row of Table 1A, preferrably on their cell surface.

[0404] Thus, the invention provides a diagnostic method useful during diagnosis of a reproductive system disorder, including cancers, which involves measuring the expression level of the gene encoding the reproductive system antigen polypeptide in tissues of the reproductive system or other cells or body fluid from an individual and comparing the measured gene expression level with a standard reproductive system antigen gene expression level, whereby an increase or decrease in the gene expression level compared to the standard is indicative of a reproductive system disorder. In specific embodiments, polynucleotides and antibodies of the invention are used to

quantitate or qualitate tissues/cells corresponding to the library source disclosed in column 7 of Table 1A expressing the corresponding reproductive system sequence disclosed in the same row of Table 1A, preferrably on their cell surface.

[0405] Where a diagnosis of a disorder in the reproductive system, including diagnosis of a tumor, has already been made according to conventional methods, the present invention is useful as a prognostic indicator, whereby patients exhibiting enhanced or depressed reproductive system antigen gene expression will experience a worse clinical outcome relative to patients expressing the gene at a level nearer the standard level.

By "assaying the expression level of the gene encoding the reproductive [0406] system associated polypeptide" is intended qualitatively or quantitatively measuring or estimating the level of the reproductive system antigen polypeptide or the level of the mRNA encoding the reproductive system antigen polypeptide in a first biological sample either directly (e.g., by determining or estimating absolute protein level or mRNA level) or relatively (e.g., by comparing to the reproductive system associated polypeptide level or mRNA level in a second biological sample). Preferably, the reproductive system antigen polypeptide expression level or mRNA level in the first biological sample is measured or estimated and compared to a standard reproductive system antigen polypeptide level or mRNA level, the standard being taken from a second biological sample obtained from an individual not having the disorder or being determined by averaging levels from a population of individuals not having a disorder of the reproductive system. As will be appreciated in the art, once a standard reproductive system antigen polypeptide level or mRNA level is known, it can be used repeatedly as a standard for comparison.

[0407] By "biological sample" is intended any biological sample obtained from an individual, cell line, tissue culture, or other source containing reproductive system antigen polypeptides (including portions thereof) or mRNA. As indicated, biological samples include body fluids (such as sera, plasma, urine, synovial fluid and spinal fluid) which contain cells expressing reproductive system antigen polypeptides, tissues of the reproductive system, and other tissue sources found to express the full length or fragments thereof of a reproductive system antigen. Methods for obtaining tissue biopsies and body fluids from mammals are well known in the art. Where the

biological sample is to include mRNA, a tissue biopsy is the preferred source.

suitable technique such as the single-step guanidinium-thiocyanate-phenol-chloroform method described in Chomczynski and Sacchi, Anal. Biochem. 162:156-159 (1987). Levels of mRNA encoding the reproductive system antigen polypeptides are then assayed using any appropriate method. These include Northern blot analysis, S1 nuclease mapping, the polymerase chain reaction (PCR), reverse transcription in combination with the polymerase chain reaction (RT-PCR), and reverse transcription in combination with the ligase chain reaction (RT-LCR).

The present invention also relates to diagnostic assays such as quantitative and diagnostic assays for detecting levels of reproductive system antigen polypeptides, in a biological sample (e.g., cells and tissues), including determination of normal and abnormal levels of polypeptides. Thus, for instance, a diagnostic assay in accordance with the invention for detecting over-expression of reproductive system antigens compared to normal control tissue samples may be used to detect the presence of tumors. Assay techniques that can be used to determine levels of a polypeptide, such as a reproductive system antigen polypeptide of the present invention in a sample derived from a host are well-known to those of skill in the art. Such assay methods include radioimmunoassays, competitive-binding assays, Western Blot analysis and ELISA assays. Assaying reproductive system antigen polypeptide levels in a biological sample can occur using any art-known method.

sample can occur using antibody-based techniques. For example, reproductive system antigen polypeptide expression in tissues can be studied with classical immunohistological methods (Jalkanen et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell . Biol. 105:3087-3096 (1987)). Other antibody-based methods useful for detecting reproductive system antigen polypeptide gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine (125 I, 121 I), carbon (14 C), sulfur (35 S), tritium (3 H), indium (112 In), and technetium (99m Tc), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

known, or suspected, to express the reproductive system related antigen gene (such as, for example, cells of the reproductive system or cancers of the reproductive system). The protein isolation methods employed herein may, for example, be such as those described in Harlow and Lane (Harlow, E. and Lane, D., 1988, "Antibodies: A Laboratory Manual", Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York), which is incorporated herein by reference in its entirety. The isolated cells can be derived from cell culture or from a patient. The analysis of cells taken from culture may be a necessary step in the assessment of cells that could be used as part of a cell-based gene therapy technique or, alternatively, to test the effect of compounds on the expression of the reproductive system related antigen gene.

[0412] For example, antibodies, or fragments of antibodies, such as those described herein, may be used to quantitatively or qualitatively detect the presence of reproductive system related antigen gene products or conserved variants or peptide fragments thereof. This can be accomplished, for example, by immunofluorescence techniques employing a fluorescently labeled antibody coupled with light microscopic, flow cytometric, or fluorimetric detection.

[0413] In a preferred embodiment, antibodies, or fragments of antibodies directed to any one or all of the predicted epitope domains of the reproductive system related antigen polypeptides (Shown in Table 1A, column 6) may be used to quantitatively or qualitatively detect the presence of reproductive system related antigen gene products or conserved variants or peptide fragments thereof. This can be accomplished, for example, by immunofluorescence techniques employing a fluorescently labeled antibody coupled with light microscopic, flow cytometric, or fluorimetric detection.

[0414] In an additional preferred embodiment, antibodies, or fragments of antibodies directed to a conformational epitope of a reproductive system related antigen may be used to quantitatively or qualitatively detect the presence of reproductive system related antigen gene products or conserved variants or peptide fragments thereof. This can be accomplished, for example, by immunofluorescence techniques employing a fluorescently labeled antibody coupled with light microscopic, flow cytometric, or fluorimetric detection.

[0415] The antibodies (or fragments thereof), and/or reproductive system related

antigen polypeptides of the present invention may, additionally, be employed histologically, as in immunofluorescence, immunoelectron microscopy or non-immunological assays, for in situ detection of reproductive system related antigen gene products or conserved variants or peptide fragments thereof. In situ detection may be accomplished by removing a histological specimen from a patient, and applying thereto a labeled antibody or reproductive system related antigen polypeptide of the present invention. The antibody (or fragment thereof) or reproductive system related antigen polypeptide is preferably applied by overlaying the labeled antibody (or fragment) onto a biological sample. Through the use of such a procedure, it is possible to determine not only the presence of the reproductive system related antigen gene product, or conserved variants or peptide fragments, or reproductive system related antigen polypeptide binding, but also its distribution in the examined tissue. Using the present invention, those of ordinary skill will readily perceive that any of a wide variety of histological methods (such as staining procedures) can be modified in order to achieve such in situ detection.

Immunoassays and non-immunoassays for reproductive system related antigen gene products or conserved variants or peptide fragments thereof will typically comprise incubating a sample, such as a biological fluid, a tissue extract, freshly harvested cells, or lysates of cells which have been incubated in cell culture, in the presence of a detectably labeled antibody capable of binding reproductive system related antigen gene products or conserved variants or peptide fragments thereof, and detecting the bound antibody by any of a number of techniques well-known in the art.

onto a solid phase support or carrier such as nitrocellulose, or other solid support which is capable of immobilizing cells, cell particles or soluble proteins. The support may then be washed with suitable buffers followed by treatment with the detectably labeled anti- reproductive system related antigen antibody or detectable reproductive system related antigen polypeptide. The solid phase support may then be washed with the buffer a second time to remove unbound antibody or polypeptide. Optionally the antibody is subsequently labeled. The amount of bound label on solid support may then be detected by conventional means.

[0418] By "solid phase support or carrier" is intended any support capable of

binding an antigen or an antibody. Well-known supports or carriers include glass, polystyrene, polypropylene, polyethylene, dextran, nylon, amylases, natural and modified celluloses, polyacrylamides, gabbros, and magnetite. The nature of the carrier can be either soluble to some extent or insoluble for the purposes of the present invention. The support material may have virtually any possible structural configuration so long as the coupled molecule is capable of binding to an antigen or antibody. Thus, the support configuration may be spherical, as in a bead, or cylindrical, as in the inside surface of a test tube, or the external surface of a rod. Alternatively, the surface may be flat such as a sheet, test strip, etc. Preferred supports include polystyrene beads. Those skilled in the art will know many other suitable carriers for binding antibody or antigen, or will be able to ascertain the same by use of routine experimentation.

[0419] The binding activity of a given lot of anti- reproductive system related antigen antibody or reproductive system related antigen polypeptide may be determined according to well known methods. Those skilled in the art will be able to determine operative and optimal assay conditions for each determination by employing routine experimentation.

levels or polynucleotide levels in a biological sample obtained from an individual, reproductive system related antigen polypeptide or polynucleotide can also be detected in vivo by imaging. For example, in one embodiment of the invention, reproductive system related antigen polypeptide and/or anti- reproductive system related antigen antibodies are used to image reproductive system diseased cells, such as neoplasms. In another embodiment, reproductive system related antigen polynucleotides of the invention (e.g., polynucleotides complementary to all or a portion of reproductive system related antigen antibodies (e.g., antibodies directed to any one or a combination of the epitopes of reproductive system related antigens, antibodies directed to a conformational epitope of reproductive system related antigens, antibodies directed to the full length polypeptide expressed on the cell surface of a mammalian cell) are used to image diseased or neoplastic cells of the reproductive system.

[0421] Antibody labels or markers for in vivo imaging of reproductive system

related antigen polypeptides include those detectable by X-radiography, NMR, MRI, CAT-scans or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma. Where in vivo imaging is used to detect enhanced levels of reproductive system related antigen polypeptides for diagnosis in humans, it may be preferable to use human antibodies or "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using techniques described herein or otherwise known in the art. For example methods for producing chimeric antibodies are known in the art. See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Tanigueti et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO 8702671; Boulianne et al., Nature 312:643'(1984); Neuberger et al., Naure 314:268 (1985).

Additionally, any reproductive system related antigen polypeptides whose presence can be detected, can be administered. For example, reproductive system [0422] related antigen perpetitives labeled with a radio-opaque or other appropriate compound can be administered and visualized in vivo, as discussed, above for labeled antibodies. Further such reproductive system related antigen polypeptides can be utilized for in vitro dagnostic procedures.

A reproductive system related antigen polypeptide-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging [0423] moiety, such as a radioisotope (for example, 131 I, 112 In, 99mTc), a radio-opaque substance, or a matrial detectable by nuclear magnetic resonance, is introduced (for example, parenteral subcutaneously or intraperitoneally) into the mammal to be examined for a disorder of the reproductive system. It will be understood in the art that the size of the stillect and the imaging system used will determine the quantity of imaging moiety needs to produce diagnostic images. In the case of a radioisotope moiety, for a humansabject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of 99mTc. The lated antibody or antibody fragment will then preferentially accurate at the location of cells which contain reproductive 998

system related antigen protein. *In vivo* tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments" (Chapter 13 in *Tumor Imaging: The Radiochemical Detection of Cancer*, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)).

With respect to antibodies, one of the ways in which the anti-reproductive [0424] system related antigen antibody can be detectably labeled is by linking the same to an enzyme and using the linked product in an enzyme immunoassay (EIA) (Voller, A., "The Enzyme Linked Immunosorbent Assay (ELISA)", 1978, Diagnostic Horizons 2:1-7, Microbiological Associates Quarterly Publication, Walkersville, MD); Voller et al., J. Clin. Pathol. 31:507-520 (1978); Butler, J.E., Meth. Enzymol. 73:482-523 (1981); Maggio, E. (ed.), 1980, Enzyme Immunoassay, CRC Press, Boca Raton, FL; Ishikawa, E. et al., (eds.), 1981, Enzyme Immunoassay, Kgaku Shoin, Tokyo). The enzyme which is bound to the antibody will react with an appropriate substrate, preferably a chromogenic substrate, in such a manner as to produce a chemical moiety which can be detected, for example, by spectrophotometric, fluorimetric or by visual means. Enzymes which can be used to detectably label the antibody include, but are not limited to, malate dehydrogenase, staphylococcal nuclease, delta-5-steroid isomerase, yeast alcohol dehydrogenase, alpha-glycerophosphate, dehydrogenase, triose phosphate isomerase, horseradish peroxidase, alkaline phosphatase, asparaginase, glucose oxidase, beta-galactosidase, ribonuclease, urease, catalase, dehydrogenase, glucoamylase glucose-6-phosphate and acetylcholinesterase. Additionally, the detection can be accomplished by colorimetric methods which employ a chromogenic substrate for the enzyme. Detection may also be accomplished by visual comparison of the extent of enzymatic reaction of a substrate in comparison with similarly prepared standards.

[0425] Detection may also be accomplished using any of a variety of other immunoassays. For example, by radioactively labeling the antibodies or antibody fragments, it is possible to detect reproductive system related antigens through the use of a radioimmunoassay (RIA) (see, for example, Weintraub, B., Principles of Radioimmunoassays, Seventh Training Course on Radioligand Assay Techniques, The Endocrine Society, March, 1986, which is incorporated by reference herein). The radioactive isotope can be detected by means including, but not limited to, a gamma

counter, a scintillation counter, or autoradiography.

[0426] It is also possible to label the antibody with a fluorescent compound. When the fluorescently labeled antibody is exposed to light of the proper wave length, its presence can then be detected due to fluorescence. Among the most commonly used fluorescent labeling compounds are fluorescein isothiocyanate, rhodamine, phycocyanin, phycocyanin, allophycocyanin, ophthaldehyde and fluorescamine.

[0427] The antibody can also be detectably labeled using fluorescence emitting metals such as ¹⁵²Eu, or others of the lanthanide series. These metals can be attached to the antibody using such metal chelating groups as diethylenetriaminepentacetic acid (DTPA) or ethylenediaminetetraacetic acid (EDTA).

[0428] The antibody also can be detectably labeled by coupling it to a chemiluminescent compound. The presence of the chemiluminescent-tagged antibody is then determined by detecting the presence of luminescence that arises during the course of a chemical reaction. Examples of particularly useful chemiluminescent labeling compounds are luminol, isoluminol, theromatic acridinium ester, imidazole, acridinium salt and oxalate ester.

[0429] Likewise, a bioluminescent compound may be used to label the antibody of the present invention. Bioluminescence is a type of chemiluminescence found in biological systems in, which a catalytic protein increases the efficiency of the chemiluminescent reaction. The presence of a bioluminescent protein is determined by detecting the presence of luminescence. Important bioluminescent compounds for purposes of labeling are luciferin, luciferase and aequorin.

Methods for Detecting Diseases of the Reproductive System, Including Cancer

[0430] In general, a disease of the reproductive system or cancer may be detected in a patient based on the presence of one or more reproductive system related antigen proteins of the invention and/or polynucleotides encoding such proteins in a biological sample (for example, blood, sera, urine, and/or tumor biopsies) obtained from the patient. In other words, such proteins and/or polynucleotides may be used as markers to indicate the presence or absence of a reproductive system disease or disorder, including cancer. Cancers that may be diagnosed, and/or prognosed using the compositions of the invention include but are not limited to, cancers of the

reproductive system. In addition, such proteins and/or polynucleotidse may be useful for the detection of other diseases and cancers, including cancers of tissues/cells corresponding to the library source disclosed in column 7 of Table 1A expressing the corresponding reproductive system related sequence disclosed in the same row of Table 1A. The binding agents provided herein generally permit detection of the level of antigen that binds to the agent in the biological sample. Polynucleotide primers and probes may be used to detect the level of mRNA encoding reproductive system related antigen polypeptides, which is also indicative of the presence or absence of a reproductive system disease or disorder, including cancer. In general, reproductive system related antigen polypeptides should be present at a level that is at least three fold higher in diseased tissue than in normal tissue.

There are a variety of assay formats known to those of ordinary skill in the art for using a binding agent to detect polypeptide markers in a sample. See, e.g., Harlow and Lane, *supra*. In general, the presence or absence of a reproductive system disease in a patient may be determined by (a) contacting a biological sample obtained from a patient with a binding agent; (b) detecting in the sample a level of polypeptide that binds to the binding agent; and (c) comparing the level of polypeptide with a predetermined cut-off value.

In a preferred embodiment, the assay involves the use of binding agent immobilized on a solid support to bind to and remove the reproductive system related antigen polypeptide of the invention from the remainder of the sample. The bound polypeptide may then be detected using a detection reagent that contains a reporter group and specifically binds to the binding agent/polypeptide complex. Such detection reagents may comprise, for example, a binding agent that specifically binds to the polypeptide or an antibody or other agent that specifically binds to the binding agent, such as an anti-immunoglobulin, protein G, protein A or a lectin. Alternatively, a competitive assay may be utilized, in which a polypeptide is labeled with a reporter group and allowed to bind to the immobilized binding agent after incubation of the binding agent with the sample. The extent to which components of the sample inhibit the binding of the labeled polypeptide to the binding agent. Suitable polypeptides for

use within such assays include reproductive system related antigen polypeptides and portions thereof, or antibodies, to which the binding agent binds, as described above.

[0433] The solid support may be any material known to those of skill in the art to which reproductive system related antigen polypeptides of the invention may be attached. For example, the solid support may be a test well in a microtiter plate or a nitrocellulose or other suitable membrane. Alternatively, the support may be a bead or disc, such as glass fiberglass, latex or a plastic material such as polystyrene or polyvinylchloride. The support may also be a magnetic particle or a fiber optic sensor, such as those disclosed, for example, in U.S. Patent No. 5,359,681. The binding agent may be immobilized on the solid support using a variety of techniques known to those of skill in the art, which are amply described in the patent and scientific literature. In the context of the present invention, the term "immobilization" refers to both noncovalent association, such as adsorption, and covalent attachment (which may be a direct linkage between the agent and functional groups on the support or may be a linkage by way of a cross-linking agent). Immobilization by adsorption to a well in a microtiter plate or to a membrane is preferred. In such cases, adsorption may be achieved by contacting the binding agent, in a suitable buffer, with the solid support for the suitable amount of time. The contact time varies with temperature, but is typically between about 1 hour and about 1 day. In general, contacting a well of plastic microtiter plate (such as polystyrene or polyvinylchloride) with an amount of binding agent ranging from about 10 ng to about 10 ug, and preferably about 100 ng to about 1 ug, is sufficient to immobilize an adequate amount of binding agent.

[0434] Covalent attachment of binding agent to a solid support may generally be achieved by first reacting the support with a bifunctional reagent that will react with both the support and a functional group, such as a hydroxyl or amino group, on the binding agent. For example, the binding agent may be covalently attached to supports having an appropriate polymer coating using benzoquinone or by condensation of an aldehyde group on the support with an amine and an active hydrogen on the binding partner (see, e.g., Pierce Immunotechnology Catalog and Handbook, 1991, at A12-A13).

Gene Therapy Methods

Also encompassed by the present invention are gene therapy methods for treating or preventing disorders, diseases and conditions. The gene therapy methods relate to the introduction of nucleic acid (DNA, RNA and antisense DNA or RNA) sequences into an animal to achieve expression of a reproductive system related antigen of the present invention. This method requires a polynucleotide which codes for a polypeptide of the present invention operatively linked to a promoter and any other genetic elements necessary for the expression of the polypeptide by the target tissue. Such gene therapy and delivery techniques are known in the art, see, for example, WO90/11092, which is herein incorporated by reference.

[0436] Thus, for example, cells from a patient may be engineered with a polynucleotide (DNA or RNA) comprising a promoter operably linked to a polynucleotide of the present invention ex vivo, with the engineered cells then being provided to a patient to be treated with the polypeptide of the present invention. Such methods are well-known in the art. For example, see Belldegrun, A., et al., J. Natl. Cancer Inst. 85: 207-216 (1993); Ferrantini, M. et al., Cancer Research 53: 1107-1112 (1993); Ferrantini, M. et al., J. Immunology 153: 4604-4615 (1994); Kaido, T., et al., Int. J. Cancer 60: 221-229 (1995); Ogura, H., et al., Cancer Research 50: 5102-5106 (1990); Santodonato, L., et al., Human Gene Therapy 7:1-10 (1996); Santodonato, L., et al., Gene Therapy 4:1246-1255 (1997); and Zhang, J.-F. et al., Cancer Gene Therapy 3: 31-38 (1996)), which are herein incorporated by reference. In one embodiment, the cells which are engineered are arterial cells. The arterial cells may be reintroduced into the patient through direct injection to the artery, the tissues surrounding the artery, or through catheter injection.

[0437] As discussed in more detail below, the polynucleotide constructs can be delivered by any method that delivers injectable materials to the cells of an animal, such as, injection into the interstitial space of tissues (heart, muscle, skin, lung, liver, and the like). The polynucleotide constructs may be delivered in a pharmaceutically acceptable liquid or aqueous carrier.

[0438] In one embodiment, the polynucleotide of the present invention is delivered as a naked polynucleotide. The term "naked" polynucleotide, DNA or RNA refers to sequences that are free from any delivery vehicle that acts to assist, promote or

facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. However, the polynucleotide of the present invention can also be delivered in liposome formulations and lipofectin formulations and the like can be prepared by methods well known to those skilled in the art. Such methods are described, for example, in U.S. Patent Nos. 5,593,972, 5,589,466, and 5,580,859, which are herein incorporated by reference.

- [0439] The polynucleotide vector constructs used in the gene therapy method are preferably constructs that will not integrate into the host genome nor will they contain sequences that allow for replication. Appropriate vectors include pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; pSVK3, pBPV, pMSG and pSVL available from Pharmacia; and pEF1/V5, pcDNA3.1, and pRc/CMV2 available from Invitrogen. Other suitable vectors will be readily apparent to the skilled artisan.
- driving the expression of the polynucleotide sequence. Suitable promoters include adenoviral promoters, such as the adenoviral major late promoter; or heterologous promoters, such as the cytomegalovirus (CMV) promoter; the respiratory syncytial virus (RSV) promoter; inducible promoters, such as the MMT promoter, the metallothionein promoter; heat shock promoters; the albumin promoter; the ApoAI promoter; human globin promoters; viral thymidine kinase promoters, such as the Herpes Simplex thymidine kinase promoter; retroviral LTRs; the b-actin promoter; and human growth hormone promoters. The promoter also may be the native promoter for the polynucleotide of the present invention.
- [0441] Unlike other gene therapy techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired polypeptide for periods of up to six months.
- [0442] The polynucleotide construct can be delivered to the interstitial space of tissues within the an animal, including of muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, blood, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, and

connective tissue. Interstitial space of the tissues comprises the intercellular, fluid, mucopolysaccharide matrix among the reticular fibers of organ tissues, elastic fibers in the walls of vessels or chambers, collagen fibers of fibrous tissues, or that same matrix within connective tissue ensheathing muscle cells or in the lacunae of bone. It is similarly the space occupied by the plasma of the circulation and the lymph fluid of the lymphatic channels. Delivery to the interstitial space of muscle tissue is preferred for the reasons discussed below. They may be conveniently delivered by injection into the tissues comprising these cells. They are preferably delivered to and expressed in persistent, non-dividing cells which are differentiated, although delivery and expression may be achieved in non-differentiated or less completely differentiated cells, such as, for example, stem cells of blood or skin fibroblasts. *In vivo* muscle cells are particularly competent in their ability to take up and express polynucleotides.

of DNA or RNA will be in the range of from about 0.05 mg/kg body weight to about 50 mg/kg body weight. Preferably the dosage will be from about 0.005 mg/kg to about 20 mg/kg and more preferably from about 0.05 mg/kg to about 5 mg/kg. Of course, as the artisan of ordinary skill will appreciate, this dosage will vary according to the tissue site of injection. The appropriate and effective dosage of nucleic acid sequence can readily be determined by those of ordinary skill in the art and may depend on the condition being treated and the route of administration.

into the interstitial space of tissues. However, other parenteral routes may also be used, such as, inhalation of an aerosol formulation particularly for delivery to lungs or bronchial tissues, throat or mucous membranes of the nose. In addition, naked DNA constructs can be delivered to arteries during angioplasty by the catheter used in the procedure.

[0445] The naked polynucleotides are delivered by any method known in the art, including, but not limited to, direct needle injection at the delivery site, intravenous injection, topical administration, catheter infusion, and so-called "gene guns". These delivery methods are known in the art.

[0446] The constructs may also be delivered with delivery vehicles such as viral sequences, viral particles, liposome formulations, lipofectin, precipitating agents, etc. Such methods of delivery are known in the art.

In certain embodiments, the polynucleotide constructs are complexed in a liposome preparation. Liposomal preparations for use in the instant invention include cationic (positively charged), anionic (negatively charged) and neutral preparations. However, cationic liposomes are particularly preferred because a tight charge complex can be formed between the cationic liposome and the polyanionic nucleic acid. Cationic liposomes have been shown to mediate intracellular delivery of plasmid DNA (Felgner et al., Proc. Natl. Acad. Sci. USA (1987) 84:7413-7416, which is herein incorporated by reference); mRNA (Malone et al., Proc. Natl. Acad. Sci. USA (1989) 86:6077-6081, which is herein incorporated by reference); and purified transcription factors (Debs et al., J. Biol. Chem. (1990) 265:10189-10192, which is herein incorporated by reference), in functional form.

[0448] Cationic liposomes are readily available. For example, N[1-2,3-dioleyloxy)propyl]-N,N,N-triethylammonium (DOTMA) liposomes particularly useful and are available under the trademark Lipofectin, from GIBCO BRL, Grand Island, N.Y., (see, also, Felgner et al., Proc. Natl Acad. Sci. USA (1987) 84:7413-7416, which is herein incorporated by reference). Other commercially available liposomes include transfectace (DDAB/DOPE) and DOTAP/DOPE (Boehringer).

Other cationic liposomes can be prepared from readily available materials using techniques well known in the art. See, e.g. PCT Publication No. WO 90/11092 (which is herein incorporated by reference) for a description of the synthesis of DOTAP (1,2-bis(oleoyloxy)-3-(trimethylammonio)propane) liposomes. Preparation of DOTMA liposomes is explained in the literature, see, e.g., P. Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417, which is herein incorporated by reference. Similar methods can be used to prepare liposomes from other cationic lipid materials.

[0450] Similarly, anionic and neutral liposomes are readily available, such as from Avanti Polar Lipids (Birmingham, Ala.), or can be easily prepared using readily available materials. Such materials include phosphatidyl choline, cholesterol, phosphatidyl ethanolamine, dioleoylphosphatidyl choline (DOPC),

dioleoylphosphatidyl glycerol (DOPG), dioleoylphoshatidyl ethanolamine (DOPE), among others. These materials can also be mixed with the DOTMA and DOTAP starting materials in appropriate ratios. Methods for making liposomes using these materials are well known in the art.

dioleoylphosphatidyl glycerol (DOPG), and dioleoylphosphatidyl ethanolamine (DOPE) can be used in various combinations to make conventional liposomes, with or without the addition of cholesterol. Thus, for example, DOPG/DOPC vesicles can be prepared by drying 50 mg each of DOPG and DOPC under a stream of nitrogen gas into a sonication vial. The sample is placed under a vacuum pump overnight and is hydrated the following day with deionized water. The sample is then sonicated for 2 hours in a capped vial, using a Heat Systems model 350 sonicator equipped with an inverted cup (bath type) probe at the maximum setting while the bath is circulated at 15EC. Alternatively, negatively charged vesicles can be prepared without sonication to produce multilamellar vesicles or by extrusion through nucleopore membranes to produce unilamellar vesicles of discrete size. Other methods are known and available to those of skill in the art.

[0452] The liposomes can comprise multilamellar vesicles (MLVs), small unilamellar vesicles (SUVs), or large unilamellar vesicles (LUVs), with SUVs being preferred. The various liposome-nucleic acid complexes are prepared using methods well known in the art. See, e.g., Straubinger et al., Methods of Immunology (1983), 101:512-527, which is herein incorporated by reference. For example, MLVs containing nucleic acid can be prepared by depositing a thin film of phospholipid on the walls of a glass tube and subsequently hydrating with a solution of the material to be encapsulated. SUVs are prepared by extended sonication of MLVs to produce a homogeneous population of unilamellar liposomes. The material to be entrapped is added to a suspension of preformed MLVs and then sonicated. When using liposomes containing cationic lipids, the dried lipid film is resuspended in an appropriate solution such as sterile water or an isotonic buffer solution such as 10 mM Tris/NaCl, sonicated, and then the preformed liposomes are mixed directly with the DNA. The liposome and DNA form a very stable complex due to binding of the positively charged liposomes to the cationic DNA. SUVs find use with small nucleic acid

fragments. LUVs are prepared by a number of methods, well known in the art. Commonly used methods include Ca²⁺-EDTA chelation (Papahadjopoulos et al., Biochim. Biophys. Acta (1975) 394:483; Wilson et al., Cell 17:77 (1979); ether injection (Deamer, D. and Bangham, A., Biochim. Biophys. Acta 443:629 (1976); Ostro et al., Biochem. Biophys. Res. Commun. 76:836 (1977); Fraley et al., Proc. Natl. Acad. Sci. USA 76:3348 (1979)); detergent dialysis (Enoch, H. and Strittmatter, P., Proc. Natl. Acad. Sci. USA 76:145 (1979)); and reverse-phase evaporation (REV) (Fraley et al., J. Biol. Chem. 255:10431 (1980); Szoka et al., Proc. Natl. Acad. Sci. USA 75:145 (1978); Schaefer-Ridder et al., Science 215:166 (1982)), which are herein incorporated by reference.

- [0453] Generally, the ratio of DNA to liposomes will be from about 10:1 to about 1:10. Preferably, the ration will be from about 5:1 to about 1:5. More preferably, the ration will be about 3:1 to about 1:3. Still more preferably, the ratio will be about 1:1.
- reports on the injection of genetic material, complexed with cationic liposomes carriers, into mice. U.S. Patent Nos. 4,897,355, 4,946,787, 5,049,386, 5,459,127, 5,589,466, 5,693,622, 5,580,859, 5,703,055, and international publication no. WO 94/9469 (which are herein incorporated by reference) provide cationic lipids for use in transfecting DNA into cells and mammals. U.S. Patent Nos. 5,589,466, 5,693,622, 5,580,859, 5,703,055, and International Publication No. WO 94/9469 provide methods for delivering DNA-cationic lipid complexes to mammals.
- [0455] In certain embodiments, cells are engineered, ex vivo or *in vivo*, using a retroviral particle containing RNA which comprises a sequence encoding a polypeptide of the present invention. Retroviruses from which the retroviral plasmid vectors may be derived include, but are not limited to, Moloney Murine Leukemia Virus, spleen necrosis virus, Rous sarcoma Virus, Harvey Sarcoma Virus, avian leukosis virus, gibbon ape leukemia virus, human immunodeficiency virus, Myeloproliferative Sarcoma Virus, and mammary tumor virus.
- [0456] The retroviral plasmid vector is employed to transduce packaging cell lines to form producer cell lines. Examples of packaging cells which may be transfected include, but are not limited to, the PE501, PA317, R-2, R-AM, PA12, T19-14X, VT-19-17-H2, RCRE, RCRIP, GP+E-86, GP+envAm12, and DAN cell lines as described

in Miller, Human Gene Therapy 1:5-14 (1990), which is incorporated herein by reference in its entirety. The vector may transduce the packaging cells through any means known in the art. Such means include, but are not limited to, electroporation. the use of liposomes, and CaPO₄ precipitation. In one alternative, the retroviral plasmid vector may be encapsulated into a liposome, or coupled to a lipid, and then administered to a host.

[0457] The producer cell line generates infectious retroviral vector particles which include polynucleotide encoding a polypeptide of the present invention. Such retroviral vector particles then may be employed, to transduce eukaryotic cells, either in vitro or in vivo. The transduced eukaryotic cells will express a polypeptide of the present invention.

In certain other embodiments, cells are engineered, ex vivo or *in vivo*, with polynucleotide contained in an adenovirus vector. Adenovirus can be manipulated such that it encodes and expresses a polypeptide of the present invention, and at the same time is inactivated in terms of its ability to replicate in a normal lytic viral life cycle. Adenovirus expression is achieved without integration of the viral DNA into the host cell chromosome, thereby alleviating concerns about insertional mutagenesis. Furthermore, adenoviruses have been used as live enteric vaccines for many years with an excellent safety profile (Schwartz, et al., Am. Rev. Respir. Dis.109:233-238 (1974)). Finally, adenovirus mediated gene transfer has been demonstrated in a number of instances including transfer of alpha-1-antitrypsin and CFTR to the lungs of cotton rats (Rosenfeld et al., Science 252:431-434 (1991); Rosenfeld et al., Cell 68:143-155 (1991)). Furthermore, extensive studies to attempt to establish adenovirus as a causative agent in human cancer were uniformly negative (Green et al., Proc. Natl. Acad. Sci. USA 76:6606 (1979)).

[0459] Suitable adenoviral vectors useful in the present invention are described, for example, in Kozarsky and Wilson, Curr. Opin. Genet. Devel. 3:499-503 (1993); Rosenfeld et al., Cell 68:143-155 (1992); Engelhardt et al., Human Genet. Ther. 4:759-769 (1993); Yang et al., Nature Genet. 7:362-369 (1994); Wilson et al., Nature 365:691-692 (1993); and U.S. Patent No. 5,652,224, which are herein incorporated by reference. For example, the adenovirus vector Ad2 is useful and can be grown in human 293 cells. These cells contain the E1 region of adenovirus and constitutively

express Ela and Elb, which complement the defective adenoviruses by providing the products of the genes deleted from the vector. In addition to Ad2, other varieties of adenovirus (e.g., Ad3, Ad5, and Ad7) are also useful in the present invention.

Preferably, the adenoviruses used in the present invention are replication deficient. Replication deficient adenoviruses require the aid of a helper virus and/or packaging cell line to form infectious particles. The resulting virus is capable of infecting cells and can express a polynucleotide of interest which is operably linked to a promoter, but cannot replicate in most cells. Replication deficient adenoviruses may be deleted in one or more of all or a portion of the following genes: E1a, E1b, E3, E4, E2a, or L1 through L5.

using an adeno-associated virus (AAV). AAVs are naturally occurring defective viruses that require helper viruses to produce infectious particles (Muzyczka, N., Curr. Topics in Microbiol. Immunol. 158:97 (1992)). It is also one of the few viruses that may integrate its DNA into non-dividing cells. Vectors containing as little as 300 base pairs of AAV can be packaged and can integrate, but space for exogenous DNA is limited to about 4.5 kb. Methods for producing and using such AAVs are known in the art. See, for example, U.S. Patent Nos. 5,139,941, 5,173,414, 5,354,678, 5,436,146, 5,474,935, 5,478,745, and 5,589,377.

will include all the sequences necessary for DNA replication, encapsidation, and host-cell integration. The polynucleotide construct is inserted into the AAV vector using standard cloning methods, such as those found in Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press (1989). The recombinant AAV vector is then transfected into packaging cells which are infected with a helper virus, using any standard technique, including lipofection, electroporation, calcium phosphate precipitation, etc. Appropriate helper viruses include adenoviruses, cytomegaloviruses, vaccinia viruses, or herpes viruses. Once the packaging cells are transfected and infected, they will produce infectious AAV viral particles which contain the polynucleotide construct. These viral particles are then used to transduce eukaryotic cells, either ex vivo or *in vivo*. The transduced cells will contain the

polynucleotide construct integrated into its genome, and will express a polypeptide of the invention...

heterologous control regions and endogenous reproductive system related antigen polynucleotide sequences (e.g., encoding a reproductive system related antigen polypeptide of the present invention) via homologous recombination (see, e.g., U.S. Patent No. 5,641,670, issued June 24, 1997; International Publication No. WO 96/29411, published September 26, 1996; International Publication No. WO 94/12650, published August 4, 1994; Koller et al., Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); and Zijlstra et al., Nature 342:435-438 (1989), which are herein incorporated by reference. This method involves the activation of a gene which is present in the target cells, but which is not normally expressed in the cells, or is expressed at a lower level than desired.

Polynucleotide constructs are made, using standard techniques known in the art, which contain the promoter with targeting sequences flanking the promoter. Suitable promoters are described herein. The targeting sequence is sufficiently complementary to an endogenous sequence to permit homologous recombination of the promoter-targeting sequence with the endogenous sequence. The targeting sequence will be sufficiently near the 5' end of the desired endogenous polynucleotide sequence so the promoter will be operably linked to the endogenous sequence upon homologous recombination.

Preferably, the amplified promoter contains distinct restriction enzyme sites on the 5' and 3' ends. Preferably, the 3' end of the first targeting sequence contains the same restriction enzyme site as the 5' end of the amplified promoter and the 5' end of the second targeting sequence contains the same restriction site as the 3' end of the amplified promoter. The amplified promoter and targeting sequences are digested and ligated together.

The promoter-targeting sequence construct is delivered to the cells, either as naked polynucleotide, or in conjunction with transfection-facilitating agents, such as liposomes, viral sequences, viral particles, whole viruses, lipofection, precipitating agents, etc., described in more detail above. The P promoter-targeting sequence can

be delivered by any method, included direct needle injection, intravenous injection, topical administration, catheter infusion, particle accelerators, etc. The methods are described in more detail below.

[0467] The promoter-targeting sequence construct is taken up by cells. Homologous recombination between the construct and the endogenous sequence takes place, such that an endogenous sequence is placed under the control of the promoter. The promoter then drives the expression of the endogenous sequence.

[0468] The polynucleotide encoding a polypeptide of the present invention may contain a secretory signal sequence that facilitates secretion of the protein. Typically, the signal sequence is positioned in the coding region of the polynucleotide to be expressed towards or at the 5' end of the coding region. The signal sequence may be homologous or heterologous to the reproductive system related antigen polynucleotide of interest and may be homologous or heterologous to the cells to be transfected. Additionally, the signal sequence may be chemically synthesized using methods known in the art.

Any mode of administration of any of the above-described polynucleotides constructs can be used so long as the mode results in the expression of one or more molecules in an amount sufficient to provide a therapeutic effect. This includes direct needle injection, systemic injection, catheter infusion, biolistic injectors, particle accelerators (i.e., "gene guns"), gelfoam sponge depots, other commercially available depot materials, osmotic pumps (e.g., Alza minipumps), oral or suppositorial solid (tablet or pill) pharmaceutical formulations, and decanting or topical applications during surgery. For example, direct injection of naked calcium phosphate-precipitated plasmid into rat liver and rat spleen or a protein-coated plasmid into the portal vein has resulted in gene expression of the foreign gene in the rat livers (Kaneda et al., Science 243:375 (1989)).

[0470] A preferred method of local administration is by direct injection. Preferably, a recombinant molecule of the present invention complexed with a delivery vehicle is administered by direct injection into or locally within the area of arteries. Administration of a composition locally within the area of arteries refers to injecting the composition centimeters and preferably, millimeters within arteries.

[0471] Another method of local administration is to contact a polynucleotide construct of the present invention in or around a surgical wound. For example, a patient can undergo surgery and the polynucleotide construct can be coated on the surface of tissue inside the wound or the construct can be injected into areas of tissue inside the wound.

Therapeutic compositions useful in systemic administration, include recombinant molecules of the present invention complexed to a targeted delivery vehicle of the present invention. Suitable delivery vehicles for use with systemic administration comprise liposomes comprising ligands for targeting the vehicle to a particular site. In specific embodiments, suitable delivery vehicles for use with systemic administration comprise liposomes comprising polypeptides of the invention for targeting the vehicle to a particular site.

injection, aerosol, oral and percutaneous (topical) delivery. Intravenous injections can be performed using methods standard in the art. Aerosol delivery can also be performed using methods standard in the art (see, for example, Stribling et al., Proc. Natl. Acad. Sci. USA 189:11277-11281, 1992, which is incorporated herein by reference). Oral delivery can be performed by complexing a polynucleotide construct of the present invention to a carrier capable of withstanding degradation by digestive enzymes in the gut of an animal. Examples of such carriers, include plastic capsules or tablets, such as those known in the art. Topical delivery can be performed by mixing a polynucleotide construct of the present invention with a lipophilic reagent (e.g., DMSO) that is capable of passing into the skin.

[0474] Determining an effective amount of substance to be delivered can depend upon a number of factors including, for example, the chemical structure and biological activity of the substance, the age and weight of the animal, the precise condition requiring treatment and its severity, and the route of administration. The frequency of treatments depends upon a number of factors, such as the amount of polynucleotide constructs administered per dose, as well as the health and history of the subject. The precise amount, number of doses, and timing of doses will be determined by the attending physician or veterinarian.

[0475] Therapeutic compositions of the present invention can be administered to any animal, preferably to mammals and birds. Preferred mammals include humans, dogs, cats, mice, rats, rabbits sheep, cattle, horses and pigs, with humans being particularly preferred.

Biological Activities

[0476] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, can be used in assays to test for one or more biological activities. If these polynucleotides or polypeptides, or agonists or antagonists of the present invention, do exhibit activity in a particular assay, it is likely that these molecules may be involved in the diseases associated with the biological activity. Thus, the polynucleotides and polypeptides, and agonists or antagonists could be used to treat, prevent diagnose and/or prognose the associated disease.

[0477] The reproductive system related antigen polynucleotides and polypeptides of the invention are predicted to have predominant expression in tissues of the reproductive system.

10478] Thus, the reproductive system related antigens of the invention may be useful as therapeutic molecules. Each would be useful for diagnosis, detection, treatment and/or prevention of diseases or disorders of the reproductive system, including, for example, injury and trauma, infections, neoplastic disorders, congenital defects, and diseases or disorders which result in infertility, complications with pregnancy, labor, or parturition, postpartum difficulties, and/or as described below in the section entitled "Reproductive System Disorders".

In a preferred embodiment, polynucleotides of the invention (e.g., a nucleic acid sequence of SEQ ID NO:X or the complement thereof; or the cDNA sequence contained in Clone ID NO:Z, or fragments or variants thereof) and/or polypeptides of the invention (e.g., an amino acid sequence contained in SEQ ID NO:Y, an amino acid sequence encoded by SEQ ID NO:X, or the complement threof, an amino acid sequence encoded by the cDNA sequence contained in Clone ID NO:Z and fragments or variants thereof as described herein) are useful for the diagnosis, detection, treatement, and/or prevention of diseases or disorders of the tissues/cells corresponding to the library source disclosed in column 7 of Table 1A expressing the

corresponding reproductive system related sequence disclosed in the same row of Table 1A.

therapeutic for cancers of the reproductive system. Treatment, diagnosis, detection, and/or prevention of disorders of the reproductive system could be carried out using a reproductive system related antigen or soluble form of a reproductive system related antigen, a reproductive system related antigen ligand, gene therapy, or ex vivo applications. Moreover, inhibitors of a reproductive system related antigen, either blocking antibodies or mutant forms, could modulate the expression of the reproductive system related antigen. These inhibitors may be useful to treat, diagnose, detect, and/or prevent diseases associated with the misregulation of a reproductive system related antigen.

In one embodiment, the invention provides a method for the specific delivery of compositions of the invention to cells (e.g., normal or diseased reproductive system cells) by administering polypeptides of the invention (e.g., reproductive system related antigen polypeptides or anti- reproductive system related antigen antibodies) that are associated with heterologous polypeptides or nucleic acids. In one example, the invention provides a method for delivering a therapeutic protein into the targeted cell (e.g., an aberrant reproductive system cell or reproductive system cancer cell). In another example, the invention provides a method for delivering a single stranded nucleic acid (e.g., antisense or ribozymes) or double stranded nucleic acid (e.g., DNA that can integrate into the cell's genome or replicate episomally and that can be transcribed) into the targeted cell.

[0482] In another embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of aberrant reproductive system cells, including, but not limited to, reproductive system tumor cells) by administering polypeptides of the invention (e.g., reproductive system related antigen polypeptides or fragments thereof, or anti- reproductive system related antigen antibodies) in association with toxins or cytotoxic prodrugs.

[0483] By "toxin" is meant compounds that bind and activate endogenous cytotoxic effector systems, radioisotopes, holotoxins, modified toxins, catalytic subunits of toxins, cytotoxins (cytotoxic agents), or any molecules or enzymes not

normally present in or on the surface of a cell that under defined conditions cause the cell's death. Toxins that may be used according to the methods of the invention include, but are not limited to, radioisotopes known in the art, compounds such as, for example, antibodies (or complement fixing containing portions thereof) that bind an inherent or induced endogenous cytotoxic effector system, thymidine kinase, endonuclease, RNAse, alpha toxin, ricin, abrin, *Pseudomonas* exotoxin A, diphtheria toxin, saporin, momordin, gelonin, pokeweed antiviral protein, alpha-sarcin and cholera toxin. "Toxin" also includes a cytostatic or cytocidal agent, a therapeutic agent or a radioactive metal ion, e.g., alpha-emitters such as, for example, ²¹³Bi, or other radioisotopes such as, for example, ¹⁰³Pd, ¹³³Xe, ¹³¹I, ⁶⁸Ge, ⁵⁷Co, ⁶⁵Zn, ⁸⁵Sr, ³²P, ³⁵S, ⁹⁰Y, ¹⁵³Sm, ¹⁵³Gd, ¹⁶⁹Yb, ⁵¹Cr, ⁵⁴Mn, ⁷⁵Se, ¹¹³Sn, ⁹⁰Yttrium, ¹¹⁷Tin, ¹⁸⁶Rhenium, ¹⁶⁶Holmium, and ¹⁸⁸Rhenium; luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin.

Techniques known in the art may be applied to label antibodies of the [0484] invention. Such techniques include, but are not limited to, the use of bifunctional conjugating agents (see e.g., U.S. Patent Nos. 5,756,065; 5,714,631; 5,696,239; 5,652,361; 5,505,931; 5,489,425; 5,435,990; 5,428,139; 5,342,604; 5,274,119; 4,994,560; and 5,808,003; the contents of each of which are hereby incorporated by reference in its entirety). A cytotoxin or cytotoxic agent includes any agent that is detrimental to cells. Examples include paclitaxol, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, tenoposide, vincristine, vinblastine, colchicin, doxorubicin, daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone, glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs or homologs thereof. Therapeutic agents include, but are not limited to, antimetabolites (e.g., methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating agents (e.g., mechlorethamine, thioepa chlorambucil, melphalan, carmustine (BSNU) (CCNU), cyclothosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cis- dichlorodiamine platinum (II) (DDP) cisplatin), anthracyclines (e.g., daunorubicin (formerly daunomycin) and doxorubicin), antibiotics (e.g., dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)), and anti-mitotic agents (e.g., vincristine and vinblastine).

By "cytotoxic prodrug" is meant a non-toxic compound that is converted by an enzyme, normally present in the cell, into a cytotoxic compound. Cytotoxic prodrugs that may be used according to the methods of the invention include, but are not limited to, glutamyl derivatives of benzoic acid mustard alkylating agent, phosphate derivatives of etoposide or mitomycin C, cytosine arabinoside, daunorubisin, and phenoxyacetamide derivatives of doxorubicin.

It will be appreciated that conditions caused by a decrease in the standard or normal level of a reproductive system related antigen activity in an individual, particularly disorders of the reproductive system, can be treated by administration of a reproductive system related antigen polypeptide (e.g., such as, for example, the complete reproductive system related antigen polypeptide, the soluble form of the extracellular domain of a reproductive system related antigen polypeptide, or cells expressing the complete protein) or agonist. Thus, the invention also provides a method of treatment of an individual in need of an increased level of reproductive system related antigen activity comprising administering to such an individual a pharmaceutical composition comprising an amount of an isolated reproductive system related antigen polypeptide of the invention, or agonist thereof (e.g., an agonistic antireproductive system related antigen antibody), effective to increase the reproductive system related antigen activity level in such an individual.

It will also be appreciated that conditions caused by a increase in the standard or normal level of reproductive system related antigen activity in an individual, particularly disorders of the reproductive system, can be treated by administration of reproductive system related antigen polypeptides (e.g., such as, for example, the complete reproductive system related antigen polypeptide, the soluble form of the extracellular domain of a reproductive system related antigen polypeptide, or cells expressing the complete protein) or antagonist (e.g., an antagonistic reproductive system related antigen antibody). Thus, the invention also provides a method of treatment of an individual in need of an decreased level of reproductive system related antigen activity comprising administering to such an individual a pharmaceutical composition comprising an amount of an isolated reproductive system related antigen polypeptide of the invention, or antagonist thereof (e.g., an antagonistic

anti- reproductive system related antigen antibody), effective to decrease the reproductive system related antigen activity level in such an individual.

[0488] In certain embodiments, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to diagnose and/or prognose diseases and/or disorders associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1A, column 7 (Tissue Distribution Library Code).

[0489] More generally, polynucleotides, translation products and antibodies corresponding to this gene may be useful for the diagnosis, prognosis, prevention, and/or treatment of diseases and/or disorders associated with the following systems.

Reproductive System Disorders

[0490] The polynucleotides or polypeptides, or agonists or antagonists of the invention may be used for the diagnosis, treatment, or prevention of diseases and/or disorders of the reproductive system. Reproductive system disorders that can be treated by the compositions of the invention, include, but are not limited to, reproductive system injuries, infections, neoplastic disorders, congenital defects, and diseases or disorders which result in infertility, complications with pregnancy, labor, or parturition, and postpartum difficulties.

disorders of the testes, including, but not limited to, testicular atrophy, testicular feminization, cryptorchism (unilateral and bilateral), anorchia, ectopic testis, epididymitis and orchitis (typically resulting from infections such as, for example, gonorrhea, mumps, tuberculosis, and syphilis), testicular torsion, vasitis nodosa, germ cell tumors (e.g., seminomas, embryonal cell carcinomas, teratocarcinomas, choriocarcinomas, yolk sac tumors, and teratomas), stromal tumors (e.g., Leydig cell tumors), hydrocele, hematocele, varicocele, spermatocele, inguinal hemia, and disorders of sperm production (e.g., immotile cilia syndrome, aspermia, asthenozoospermia, azoospermia, oligospermia, and teratozoospermia).

[0492] Reproductive system disorders also include, but are not limited to, disorders of the prostate gland, such as acute non-bacterial prostatitis, chronic non-bacterial prostatitis, acute bacterial prostatitis, chronic bacterial prostatitis, prostatodystonia, prostatosis, granulomatous prostatitis, malacoplakia, benign prostatic hypertrophy or hyperplasia, and prostate neoplastic disorders, including adenocarcinomas, transitional cell carcinomas, ductal carcinomas, and squamous cell carcinomas.

[0493] Additionally, the compositions of the invention may be useful in the diagnosis, treatment, and/or prevention of disorders or diseases of the penis and urethra, including, but not limited to, inflammatory disorders, such as balanoposthitis, balanitis xerotica obliterans, phimosis, paraphimosis, syphilis, herpes simplex virus, gonorrhea, non-gonococcal urethritis, chlamydia, mycoplasma, trichomonas, HIV, AIDS, Reiter's syndrome, condyloma acuminatum, condyloma latum, and pearly penile papules; urethral abnormalities, such as hypospadias, epispadias, and phimosis; premalignant lesions, including Erythroplasia of Queyrat, Bowen's disease, Bowenoid paplosis, giant condyloma of Buscke-Lowenstein, and varrucous carcinoma; penile cancers, including squamous cell carcinomas, carcinoma in situ, verrucous carcinoma, and disseminated penile carcinoma; urethral neoplastic disorders, including penile urethral carcinoma, bulbomembranous urethral carcinoma, and prostatic urethral carcinoma; and erectile disorders, such as priapism, Peyronie's disease, erectile dysfunction, and impotence.

limited to, vasculititis and CBAVD (congenital bilateral absence of the vas deferens); additionally, the polynucleotides, polypeptides, and agonists or antagonists of the present invention may be used in the diagnosis, treatment, and/or prevention of diseases and/or disorders of the seminal vesicles, including but not limited to, hydatid disease, congenital chloride diarrhea, and polycystic kidney disease.

[0495] Other disorders and/or diseases of the male reproductive system that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, Klinefelter's syndrome, Young's syndrome, premature ejaculation, diabetes mellitus, cystic fibrosis, Kartagener's syndrome, high fever, multiple sclerosis, and gynecomastia.

further, the polynucleotides, polypeptides, and agonists or antagonists of the present invention may be used in the diagnosis, treatment, and/or prevention of diseases and/or disorders of the vagina and vulva, including, but not limited to, bacterial vaginosis, candida vaginitis, herpes simplex virus, chancroid, granuloma inguinale, lymphogranuloma venereum, scabies, human papillomavirus, vaginal trauma, vulvar trauma, adenosis, chlamydia vaginitis, gonorrhea, trichomonas vaginitis, condyloma acuminatum, syphilis, molluscum contagiosum, atrophic vaginitis, Paget's disease, lichen sclerosus, lichen planus, vulvodynia, toxic shock syndrome, vaginismus, vulvovaginitis, vulvar vestibulitis, and neoplastic disorders, such as squamous cell hyperplasia, clear cell carcinoma, basal cell carcinoma, melanomas, cancer of Bartholin's gland, and vulvar intraepithelial neoplasia.

Disorders and/or diseases of the uterus that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, dysmenorrhea, retroverted uterus, endometriosis, fibroids, adenomyosis, anovulatory bleeding, amenorrhea, Cushing's syndrome, hydatidiform moles, Asherman's syndrome, premature menopause, precocious puberty, uterine polyps, dysfunctional uterine bleeding (e.g., due to aberrant hormonal signals), and neoplastic disorders, such as adenocarcinomas, keiomyosarcomas, and sarcomas. Additionally, the polypeptides, polynucleotides, or agonists or antagonists of the invention may be useful as a marker or detector of, as well as in the diagnosis, treatment, and/or prevention of congenital uterine abnormalities, such as bicornuate uterus, septate uterus, simple unicornuate uterus, unicornuate uterus with a noncavitary rudimentary horn, unicornuate uterus with a non-communicating cavitary rudimentary horn, unicornuate uterus with a communicating cavitary rudimentary horn, unicornuate uterus with a communicating cavitary horn, arcuate uterus, uterine didelfus, and T-shaped uterus.

[0498] Ovarian diseases and/or disorders that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, anovulation, polycystic ovary syndrome (Stein-Leventhal syndrome), ovarian cysts, ovarian hypofunction, ovarian insensitivity to gonadotropins, ovarian overproduction of androgens, right ovarian vein syndrome, amenorrhea, hirutism, and ovarian cancer (including, but not limited to, primary and secondary cancerous growth, Sertoli-Leydig tumors, endometriod carcinoma of the ovary, ovarian papillary serous

adenocarcinoma, ovarian mucinous adenocarcinoma, and Ovarian Krukenberg tumors).

[0499] Cervical diseases and/or disorders that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, cervicitis, chronic cervicitis, mucopurulent cervicitis, cervical dysplasia, cervical polyps, Nabothian cysts, cervical erosion, cervical incompetence, and cervical neoplasms (including, for example, cervical carcinoma, squamous metaplasia, squamous cell carcinoma, adenosquamous cell neoplasia, and columnar cell neoplasia).

[0500] Additionally, diseases and/or disorders of the reproductive system that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, disorders and/or diseases of pregnancy, including miscarriage and stillbirth, such as early abortion, late abortion, spontaneous abortion, induced abortion, therapeutic abortion, threatened abortion, missed abortion, incomplete abortion, complete abortion, habitual abortion, missed abortion, and septic abortion; ectopic pregnancy, anemia, Rh incompatibility, vaginal bleeding during pregnancy, gestational diabetes, intrauterine growth retardation, polyhydramnios, HELLP syndrome, abruptio placentae, placenta previa, hyperemesis, preeclampsia, eclampsia, herpes gestationis, and urticaria of pregnancy. Additionally, the polynucleotides, polypeptides, and agonists or antagonists of the present invention may be used in the diagnosis, treatment, and/or prevention of diseases that can complicate pregnancy, including heart disease, heart failure, rheumatic heart disease, congenital heart disease, mitral valve prolapse, high blood pressure, anemia, kidney disease, infectious disease (e.g., rubella, cytomegalovirus, toxoplasmosis, infectious hepatitis, chlamydia, HIV, AIDS, and genital herpes), diabetes mellitus, Graves' disease, thyroiditis, hypothyroidism, Hashimoto's thyroiditis, chronic active hepatitis, cirrhosis of the liver, primary biliary cirrhosis, asthma, systemic lupus eryematosis, rheumatoid arthritis, myasthenia gravis, idiopathic thrombocytopenic purpura, appendicitis, ovarian cysts, gallbladder disorders, and obstruction of the intestine.

[0501] Complications associated with labor and parturition that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, premature rupture of the membranes, pre-term labor, post-term pregnancy,

postmaturity, labor that progresses too slowly, fetal distress (e.g., abnormal heart rate (fetal or maternal), breathing problems, and abnormal fetal position), shoulder dystocia, prolapsed umbilical cord, amniotic fluid embolism, and aberrant uterine bleeding.

[0502] Further, diseases and/or disorders of the postdelivery period, that may be diagnosed, treated, and/or prevented with the compositions of the invention, include, but are not limited to, endometritis, myometritis, parametritis, peritonitis, pelvic thrombophlebitis, pulmonary embolism, endotoxemia, pyelonephritis, saphenous thrombophlebitis, mastitis, cystitis, postpartum hemorrhage, and inverted uterus.

[0503] Other disorders and/or diseases of the female reproductive system that may be diagnosed, treated, and/or prevented by the polynucleotides, polypeptides, and agonists or antagonists of the present invention include, but are not limited to, Turner's syndrome, pseudohermaphroditism, premenstrual syndrome, pelvic inflammatory disease, pelvic congestion (vascular engorgement), frigidity, anorgasmia, dyspareunia, ruptured fallopian tube, and Mittelschmerz.

Immune Activity

Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, diagnosing and/or prognosing diseases, disorders, and/or conditions of the immune system, by, for example, activating or inhibiting the proliferation, differentiation, or mobilization (chemotaxis) of immune cells. Immune cells develop through a process called hematopoiesis, producing myeloid (platelets, red blood cells, neutrophils, and macrophages) and lymphoid (B and T lymphocytes) cells from pluripotent stem cells. The etiology of these immune diseases, disorders, and/or conditions may be genetic, somatic, such as cancer and some autoimmune diseases, acquired (e.g., by chemotherapy or toxins), or infectious. Moreover, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention can be used as a marker or detector of a particular immune system disease or disorder.

[0505] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to treat diseases and disorders of the immune system and/or to inhibit or enhance an

immune response generated by cells associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1A, column 7 (Tissue Distribution Library Code).

105061 Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, diagnosing, and/or immunodeficiencies, prognosing including both congenital and acquired immunodeficiencies. Examples of B cell immunodeficiencies in which immunoglobulin levels B cell function and/or B cell numbers are decreased include: X-linked agammaglobulinemia (Bruton's disease), X-linked infantile agammaglobulinemia, X-linked immunodeficiency with hyper IgM, non X-linked immunodeficiency with hyper IgM, X-linked lymphoproliferative syndrome (XLP), agammaglobulinemia including congenital and acquired agammaglobulinemia, adult onset agammaglobulinemia, late-onset agammaglobulinemia, dysgammaglobulinemia, hypogammaglobulinemia, unspecified hypogammaglobulinemia, recessive agammaglobulinemia (Swiss type), Selective IgM deficiency, selective IgA deficiency, selective IgG subclass deficiencies, IgG subclass deficiency (with or without IgA deficiency), Ig deficiency with increased IgM, IgG and IgA deficiency with increased IgM, antibody deficiency with normal or elevated Igs, Ig heavy chain deletions, kappa chain deficiency, B cell lymphoproliferative disorder (BLPD), common variable immunodeficiency (CVID), common variable immunodeficiency (CVI) (acquired), and transient hypogammaglobulinemia of infancy.

[0507] In specific embodiments, ataxia-telangiectasia or conditions associated with ataxia-telangiectasia are treated, prevented, diagnosed, and/or prognosing using the polypeptides or polynucleotides of the invention, and/or agonists or antagonists thereof.

[0508] Examples of congenital immunodeficiencies in which T cell and/or B cell function and/or number is decreased include, but are not limited to: DiGeorge anomaly, severe combined immunodeficiencies (SCID) (including, but not limited to, X-linked SCID, autosomal recessive SCID, adenosine deaminase deficiency, purine nucleoside phosphorylase (PNP) deficiency, Class II MHC deficiency (Bare lymphocyte syndrome), Wiskott-Aldrich syndrome, and ataxia telangiectasia), thymic hypoplasia, third and fourth pharyngeal pouch syndrome, 22q11.2 deletion, chronic

mucocutaneous candidiasis, natural killer cell deficiency (NK), idiopathic CD4+ T-lymphocytopenia, immunodeficiency with predominant T cell defect (unspecified), and unspecified immunodeficiency of cell mediated immunity.

- [0509] In specific embodiments, DiGeorge anomaly or conditions associated with DiGeorge anomaly are treated, prevented, diagnosed, and/or prognosed using polypeptides or polynucleotides of the invention, or antagonists or agonists thereof.
- [0510] Other immunodeficiencies that may be treated, prevented, diagnosed, and/or prognosed using polypeptides or polynucleotides of the invention, and/or agonists or antagonists thereof, include, but are not limited to, chronic granulomatous disease, Chédiak-Higashi syndrome, myeloperoxidase deficiency, leukocyte glucose-6-phosphate dehydrogenase deficiency, X-linked lymphoproliferative syndrome (XLP), leukocyte adhesion deficiency, complement component deficiencies (including C1, C2, C3, C4, C5, C6, C7, C8 and/or C9 deficiencies), reticular dysgenesis, thymic alymphoplasia-aplasia, immunodeficiency with thymoma, severe congenital leukopenia, dysplasia with immunodeficiency, neonatal neutropenia, short limbed dwarfism, and Nezelof syndrome-combined immunodeficiency with Igs.
- [0511] In a preferred embodiment, the immunodeficiencies and/or conditions associated with the immunodeficiencies recited above are treated, prevented, diagnosed and/or prognosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.
- [0512] In a preferred embodiment polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used as an agent to boost immunoresponsiveness among immunodeficient individuals. In specific embodiments, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used as an agent to boost immunoresponsiveness among B cell and/or T cell immunodeficient individuals.
- [0513] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, diagnosing and/or prognosing autoimmune disorders. Many autoimmune disorders result from inappropriate recognition of self as foreign material by immune cells. This inappropriate recognition results in an immune response leading to the destruction of the host tissue. Therefore, the administration of polynucleotides and polypeptides of

the invention that can inhibit an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing autoimmune disorders.

Autoimmune diseases or disorders that may be treated, prevented, [0514] diagnosed and/or prognosed by polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include, but are not limited to, one or more of the following: systemic lupus erythematosus, rheumatoid arthritis, ankylosing spondylitis, multiple sclerosis, autoimmune thyroiditis, Hashimoto's thyroiditis, autoimmune hemolytic anemia, hemolytic anemia, thrombocytopenia, autoimmune thrombocytopenia purpura, autoimmune neonatal thrombocytopenia, idiopathic thrombocytopenia purpura, purpura (e.g., Henloch-Scoenlein purpura), autoimmunocytopenia, Goodpasture's syndrome, Pemphigus vulgaris, myasthenia gravis, Grave's disease (hyperthyroidism), and insulin-resistant diabetes mellitus.

[0515] Additional disorders that are likely to have an autoimmune component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis. myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disorders.

[0516] Additional disorders that are likely to have an autoimmune component that may be treated, prevented, diagnosed and/or prognosed with the compositions of the invention include, but are not limited to, scleroderma with anti-collagen antibodies (often characterized, e.g., by nucleolar and other nuclear antibodies), mixed connective tissue disease (often characterized, e.g., by antibodies to extractable nuclear antigens (e.g., ribonucleoprotein)), polymyositis (often characterized, e.g., by nonhistone ANA), pernicious anemia (often characterized, e.g., by antiparietal cell, microsomes, and intrinsic factor antibodies), idiopathic Addison's disease (often characterized, e.g., by humoral and cell-mediated adrenal cytotoxicity, infertility (often characterized, e.g., by antispermatozoal antibodies), glomerulonephritis (often characterized, e.g., by glomerular basement membrane antibodies or immune complexes), bullous

pemphigoid (often characterized, e.g., by IgG and complement in basement membrane), Sjogren's syndrome (often characterized, e.g., by multiple tissue antibodies, and/or a specific nonhistone ANA (SS-B)), diabetes mellitus (often characterized, e.g., by cell-mediated and humoral islet cell antibodies), and adrenergic drug resistance (including adrenergic drug resistance with asthma or cystic fibrosis) (often characterized, e.g., by beta-adrenergic receptor antibodies).

treated, prevented, diagnosed and/or prognosed with the compositions of the invention include, but are not limited to, chronic active hepatitis (often characterized, e.g., by smooth muscle antibodies), primary biliary cirrhosis (often characterized, e.g., by mitochondria antibodies), other endocrine gland failure (often characterized, e.g., by specific tissue antibodies in some cases), vitiligo (often characterized, e.g., by melanocyte antibodies), vasculitis (often characterized, e.g., by Ig and complement in vessel walls and/or low serum complement), post-MI (often characterized, e.g., by myocardial antibodies), cardiotomy syndrome (often characterized, e.g., by myocardial antibodies), urticaria (often characterized, e.g., by IgG and IgM antibodies to IgE), atopic dermatitis (often characterized, e.g., by IgG and IgM antibodies to IgE), asthma (often characterized, e.g., by IgG and IgM antibodies to IgE), asthma (often characterized, e.g., by IgG and IgM antibodies to IgE), and many other inflammatory, granulomatous, degenerative, and atrophic disorders.

[0518] In a preferred embodiment, the autoimmune diseases and disorders and/or conditions associated with the diseases and disorders recited above are treated, prevented, diagnosed and/or prognosed using for example, antagonists or agonists, polypeptides or polynucleotides, or antibodies of the present invention. In a specific preferred embodiment, rheumatoid arthritis is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0519] In another specific preferred embodiment, systemic lupus erythematosus is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention. In another specific preferred embodiment, idiopathic thrombocytopenia purpura is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0520] In another specific preferred embodiment IgA nephropathy is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0521] In a preferred embodiment, the autoimmune diseases and disorders and/or conditions associated with the diseases and disorders recited above are treated, prevented, diagnosed and/or prognosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0522] In preferred embodiments, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a immunosuppressive agent(s).

Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of [0523] the present invention may be useful in treating, preventing, prognosing, and/or diseases, disorders, and/or conditions of hematopoietic cells. diagnosing Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat or prevent those diseases, disorders, and/or conditions associated with a decrease in certain (or many) types hematopoietic cells, including but not limited to, leukopenia, neutropenia, anemia, and thrombocytopenia. Alternatively, Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat or prevent those diseases, disorders, and/or conditions associated with an increase in certain (or many) types of hematopoietic cells, including but not limited to, histiocytosis.

[0524] Allergic reactions and conditions, such as asthma (particularly allergic asthma) or other respiratory problems, may also be treated, prevented, diagnosed and/or prognosed using polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof. Moreover, these molecules can be used to treat, prevent, prognose, and/or diagnose anaphylaxis, hypersensitivity to an antigenic molecule, or blood group incompatibility.

[0525] Additionally, polypeptides or polynucleotides of the invention, and/or agonists or antagonists thereof, may be used to treat, prevent, diagnose and/or

prognose IgE-mediated allergic reactions. Such allergic reactions include, but are not limited to, asthma, rhinitis, and eczema. In specific embodiments, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to modulate IgE concentrations in vitro or in vivo.

105261 Moreover, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention have uses in the diagnosis, prognosis, prevention, and/or treatment of inflammatory conditions. For example, since polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists of the invention may inhibit the activation, proliferation and/or differentiation of cells involved in an inflammatory response, these molecules can be used to prevent and/or treat chronic and acute inflammatory conditions. Such inflammatory conditions include, but are not limited to, for example, inflammation associated with infection (e.g., septic shock, sepsis, or systemic inflammatory response syndrome), ischemiareperfusion injury, endotoxin lethality, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine induced lung injury, inflammatory bowel disease, Crohn's disease, over production of cytokines (e.g., TNF or IL-1.), respiratory disorders (e.g., asthma and allergy); gastrointestinal disorders (e.g., inflammatory bowel disease); cancers (e.g., gastric, ovarian, lung, bladder, liver, and breast); CNS disorders (e.g., multiple sclerosis; ischemic brain injury and/or stroke, traumatic brain injury, neurodegenerative disorders (e.g., Parkinson's disease and Alzheimer's disease); AIDS-related dementia; and prion disease); cardiovascular disorders (e.g., atherosclerosis, myocarditis, cardiovascular disease, and cardiopulmonary bypass complications); as well as many additional diseases, conditions, and disorders that are characterized by inflammation (e.g., hepatitis, rheumatoid arthritis, gout, trauma, pancreatitis, sarcoidosis, dermatitis, renal ischemia-reperfusion injury, Grave's disease, systemic lupus erythematosus, diabetes mellitus, and allogenic transplant rejection).

[0527] Because inflammation is a fundamental defense mechanism, inflammatory disorders can effect virtually any tissue of the body. Accordingly, polynucleotides, polypeptides, and antibodies of the invention, as well as agonists or antagonists thereof, have uses in the treatment of tissue-specific inflammatory disorders, including, but not limited to, adrenalitis, alveolitis, angiocholecystitis, appendicitis,

balanitis, blepharitis, bronchitis, bursitis, carditis, cellulitis, cervicitis, cholecystitis, chorditis, cochlitis, colitis, conjunctivitis, cystitis, dermatitis, diverticulitis, encephalitis, endocarditis, esophagitis, eustachitis, fibrositis, folliculitis, gastritis, gastroenteritis, gingivitis, glossitis, hepatosplenitis, keratitis, labyrinthitis, laryngitis, lymphangitis, mastitis, media otitis, meningitis, metritis, mucitis, myocarditis, myosititis, myringitis, nephritis, neuritis, orchitis, osteochondritis, otitis, pericarditis, peritendonitis, peritonitis, pharyngitis, phlebitis, poliomyelitis, prostatitis, pulpitis, retinitis, rhinitis, salpingitis, scleritis, sclerochoroiditis, scrotitis, sinusitis, spondylitis, steatitis, stomatitis, synovitis, syringitis, tendonitis, tonsillitis, urethritis, and vaginitis.

In specific embodiments, polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, are useful to diagnose, prognose, prevent, and/or treat organ transplant rejections and graft-versus-host disease. Organ rejection occurs by host immune cell destruction of the transplanted tissue through an immune response. Similarly, an immune response is also involved in GVHD, but, in this case, the foreign transplanted immune cells destroy the host tissues. Polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, that inhibit an immune response, particularly the activation, proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing organ rejection or GVHD. In specific embodiments, polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, that inhibit an immune response, particularly the activation, proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing experimental allergic and hyperacute xenograft rejection.

[0529] In other embodiments, polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, are useful to diagnose, prognose, prevent, and/or treat immune complex diseases, including, but not limited to, serum sickness, post streptococcal glomerulonephritis, polyarteritis nodosa, and immune complex-induced vasculitis.

[0530] Polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the invention can be used to treat, detect, and/or prevent infectious agents. For example, by increasing the immune response, particularly increasing the proliferation activation and/or differentiation of B and/or T cells, infectious diseases may be treated,

detected, and/or prevented. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may also directly inhibit the infectious agent (refer to section of application listing infectious agents, etc), without necessarily eliciting an immune response.

[0531] In another embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a vaccine adjuvant that enhances immune responsiveness to an antigen. In a specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance tumor-specific immune responses.

In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance anti-viral immune responses. Anti-viral immune responses that may be enhanced using the compositions of the invention as an adjuvant, include virus and virus associated diseases or symptoms described herein or otherwise known in the art. In specific embodiments, the compositions of the invention are used as an adjuvant to enhance an immune response to a virus, disease, or symptom selected from the group consisting of: AIDS, meningitis, Dengue, EBV, and hepatitis (e.g., hepatitis B). In another specific embodiment, the compositions of the invention are used as an adjuvant to enhance an immune response to a virus, disease, or symptom selected from the group consisting of: HIV/AIDS, respiratory syncytial virus, Dengue, rotavirus, Japanese B encephalitis, influenza A and B, parainfluenza, measles, cytomegalovirus, rabies, Junin, Chikungunya, Rift Valley Fever, herpes simplex, and yellow fever.

[0533] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance anti-bacterial or anti-fungal immune responses. Anti-bacterial or anti-fungal immune responses that may be enhanced using the compositions of the invention as an adjuvant, include bacteria or fungus and bacteria or fungus associated diseases or symptoms described herein or otherwise known in the art. In specific embodiments, the compositions of the invention are used as an adjuvant to enhance an immune

response to a bacteria or fungus, disease, or symptom selected from the group consisting of: tetanus, Diphtheria, botulism, and meningitis type B.

[0534] In another specific embodiment, the compositions of the invention are used as an adjuvant to enhance an immune response to a bacteria or fungus, disease, or symptom selected from the group consisting of: Vibrio cholerae, Mycobacterium leprae, Salmonella typhi, Salmonella paratyphi, Meisseria meningitidis, Streptococcus pneumoniae, Group B streptococcus, Shigella spp., Enterotoxigenic Escherichia coli, Enterohemorrhagic E. coli, and Borrelia burgdorferi.

In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance anti-parasitic immune responses. Anti-parasitic immune responses that may be enhanced using the compositions of the invention as an adjuvant, include parasite and parasite associated diseases or symptoms described herein or otherwise known in the art. In specific embodiments, the compositions of the invention are used as an adjuvant to enhance an immune response to a parasite. In another specific embodiment, the compositions of the invention are used as an adjuvant to enhance an immune response to Plasmodium (malaria) or Leishmania.

[0536] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may also be employed to treat infectious diseases including silicosis, sarcoidosis, and idiopathic pulmonary fibrosis; for example, by preventing the recruitment and activation of mononuclear phagocytes.

[0537] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an antigen for the generation of antibodies to inhibit or enhance immune mediated responses against polypeptides of the invention.

[0538] In one embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are administered to an animal (e.g., mouse, rat, rabbit, hamster, guinea pig, pigs, micro-pig, chicken, camel, goat, horse, cow, sheep, dog, cat, non-human primate, and human, most preferably human) to boost the immune system to produce increased quantities of one or more antibodies (e.g., IgG, IgA, IgM, and IgE), to induce higher affinity antibody production and

immunoglobulin class switching (e.g., IgG, IgA, IgM, and IgE), and/or to increase an immune response.

- [0539] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a stimulator of B cell responsiveness to pathogens.
- [0540] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an activator of T cells.
- [0541] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent that elevates the immune status of an individual prior to their receipt of immunosuppressive therapies.
- [0542] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to induce higher affinity antibodies.
- [0543] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to increase serum immunoglobulin concentrations.
- [0544] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to accelerate recovery of immunocompromised individuals.
- [0545] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to boost immunoresponsiveness among aged populations and/or neonates.
- In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an immune system enhancer prior to, during, or after bone marrow transplant and/or other transplants (e.g., allogeneic or xenogeneic organ transplantation). With respect to transplantation, compositions of the invention may be administered prior to, concomitant with, and/or after transplantation. In a specific embodiment, compositions of the invention are administered after transplantation, prior to the beginning of recovery of T-cell populations. In another specific embodiment, compositions of the invention are first

administered after transplantation after the beginning of recovery of T cell populations, but prior to full recovery of B cell populations.

[0547] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to boost immunoresponsiveness among individuals having an acquired loss of B cell function. Conditions resulting in an acquired loss of B cell function that may be ameliorated or treated by administering the polypeptides, antibodies, polynucleotides and/or agonists or antagonists thereof, include, but are not limited to, HIV Infection, AIDS, bone marrow transplant, and B cell chronic lymphocytic leukemia (CLL).

In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to boost immunoresponsiveness among individuals having a temporary immune deficiency. Conditions resulting in a temporary immune deficiency that may be ameliorated or treated by administering the polypeptides, antibodies, polynucleotides and/or agonists or antagonists thereof, include, but are not limited to, recovery from viral infections (e.g., influenza), conditions associated with malnutrition, recovery from infectious mononucleosis, or conditions associated with stress, recovery from measles, recovery from blood transfusion, and recovery from surgery.

In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a regulator of antigen presentation by monocytes, dendritic cells, and/or B-cells. In one embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention enhance antigen presentation or antagonizes antigen presentation in vitro or in vivo. Moreover, in related embodiments, said enhancement or antagonism of antigen presentation may be useful as an anti-tumor treatment or to modulate the immune system.

[0550] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to direct an individual's immune system towards development of a humoral response (i.e. TH2) as opposed to a TH1 cellular response.

[0551] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means to induce

tumor proliferation and thus make it more susceptible to anti-neoplastic agents. For example, multiple myeloma is a slowly dividing disease and is thus refractory to virtually all anti-neoplastic regimens. If these cells were forced to proliferate more rapidly their susceptibility profile would likely change.

- [0552] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a stimulator of B cell production in pathologies such as AIDS, chronic lymphocyte disorder and/or Common Variable Immunodificiency.
- [0553] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for generation and/or regeneration of lymphoid tissues following surgery, trauma or genetic defect. In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used in the pretreatment of bone marrow samples prior to transplant.
- [0554] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a gene-based therapy for genetically inherited disorders resulting in immuno-incompetence/immunodeficiency such as observed among SCID patients.
- [0555] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of activating monocytes/macrophages to defend against parasitic diseases that effect monocytes such as Leishmania.
- [0556] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of regulating secreted cytokines that are elicited by polypeptides of the invention.
- [0557] In another embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used in one or more of the applications decribed herein, as they may apply to veterinary medicine.
- [0558] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of blocking various aspects of immune responses to foreign agents or self. Examples of diseases or conditions in which blocking of certain aspects of immune responses may be

desired include autoimmune disorders such as lupus, and arthritis, as well as immunoresponsiveness to skin allergies, inflammation, bowel disease, injury and diseases/disorders associated with pathogens.

[0559] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for preventing the B cell proliferation and Ig secretion associated with autoimmune diseases such as idiopathic thrombocytopenic purpura, systemic lupus erythematosus and multiple sclerosis.

[0560] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a inhibitor of B and/or T cell migration in endothelial cells. This activity disrupts tissue architecture or cognate responses and is useful, for example in disrupting immune responses, and blocking sepsis.

[0561] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for chronic hypergammaglobulinemia evident in such diseases as monoclonal gammopathy of undetermined significance (MGUS), Waldenstrom's disease, related idiopathic monoclonal gammopathies, and plasmacytomas.

[0562] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be employed for instance to inhibit polypeptide chemotaxis and activation of macrophages and their precursors, and of neutrophils, basophils, B lymphocytes and some T-cell subsets, e.g., activated and CD8 cytotoxic T cells and natural killer cells, in certain autoimmune and chronic inflammatory and infective diseases. Examples of autoimmune diseases are described herein and include multiple sclerosis, and insulin-dependent diabetes.

[0563] The polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may also be employed to treat idiopathic hypereosinophilic syndrome by, for example, preventing eosinophil production and migration.

[0564] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used to enhance or inhibit complement mediated cell lysis.

[0565] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used to enhance or inhibit antibody dependent cellular cytotoxicity.

[0566] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may also be employed for treating atherosclerosis, for example, by preventing monocyte infiltration in the artery wall.

[0567] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be employed to treat adult respiratory distress syndrome (ARDS).

[0568] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be useful for stimulating wound and tissue repair, stimulating angiogenesis, and/or stimulating the repair of vascular or lymphatic diseases or disorders. Additionally, agonists and antagonists of the invention may be used to stimulate the regeneration of mucosal surfaces.

[0569] In a specific embodiment, polynucleotides or polypeptides, and/or agonists thereof are used to diagnose, prognose, treat, and/or prevent a disorder characterized by primary or acquired immunodeficiency, deficient serum immunoglobulin production, recurrent infections, and/or immune system dysfunction. polynucleotides or polypeptides, and/or agonists thereof may be used to treat or prevent infections of the joints, bones, skin, and/or parotid glands, blood-borne infections (e.g., sepsis, meningitis, septic arthritis, and/or osteomyelitis), autoimmune diseases (e.g., those disclosed herein), inflammatory disorders, and malignancies, and/or any disease or disorder or condition associated with these infections, diseases, disorders and/or malignancies) including, but not limited to, CVID, other primary immune deficiencies, HIV disease, CLL, recurrent bronchitis, sinusitis, otitis media, conjunctivitis, pneumonia, hepatitis, meningitis, herpes zoster (e.g., severe herpes zoster), and/or pneumocystis carnii. Other diseases and disorders that may be prevented, diagnosed, prognosed, and/or treated with polynucleotides or polypeptides, and/or agonists of the present invention include, but are not limited to, HIV infection, HTLV-BLV infection, lymphopenia, phagocyte bactericidal dysfunction anemia, thrombocytopenia, and hemoglobinuria.

[0570] In another embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention are used to treat, and/or diagnose an individual having common variable immunodeficiency disease ("CVID"; also known as "acquired agammaglobulinemia" and "acquired hypogammaglobulinemia") or a subset of this disease.

- In a specific embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to diagnose, prognose, prevent, and/or treat cancers or neoplasms including immune cell or immune tissue-related cancers or neoplasms. Examples of cancers or neoplasms that may be prevented, diagnosed, or treated by polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include, but are not limited to, acute myelogenous leukemia, chronic myelogenous leukemia, Hodgkin's disease, non-Hodgkin's lymphoma, acute lymphocytic anemia (ALL) Chronic lymphocyte leukemia, plasmacytomas, multiple myeloma, Burkitt's lymphoma, EBV-transformed diseases, and/or diseases and disorders described in the section entitled "Hyperproliferative Disorders" elsewhere herein.
- [0572] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for decreasing cellular proliferation of Large B-cell Lymphomas.
- [0573] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of decreasing the involvement of B cells and Ig associated with Chronic Myelogenous Leukemia.
- [0574] In specific embodiments, the compositions of the invention are used as an agent to boost immunoresponsiveness among B cell immunodeficient individuals, such as, for example, an individual who has undergone a partial or complete splenectomy.
- [0575] Antagonists of the invention include, for example, binding and/or inhibitory antibodies, antisense nucleic acids, ribozymes or soluble forms of the polypeptides of the present invention (e.g., Fc fusion protein; see, e.g., Example 9). Agonists of the invention include, for example, binding or stimulatory antibodies, and soluble forms of the polypeptides (e.g., Fc fusion proteins; see, e.g., Example 9).

Polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be employed in a composition with a pharmaceutically acceptable carrier, e.g., as described herein.

In another embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are administered to an animal (including, but not limited to, those listed above, and also including transgenic animals) incapable of producing functional endogenous antibody molecules or having an otherwise compromised endogenous immune system, but which is capable of producing human immunoglobulin molecules by means of a reconstituted or partially reconstituted immune system from another animal (see, e.g., published PCT Application Nos. WO98/24893, WO/9634096, WO/9633735, and WO/9110741). Administration of polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention to such animals is useful for the generation of monoclonal antibodies against the polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention.

Blood-Related Disorders

antagonists of the present invention may be used to modulate hemostatic (the stopping of bleeding) or thrombolytic (clot dissolving) activity. For example, by increasing hemostatic or thrombolytic activity, polynucleotides or polypeptides, and/or agonists or antagonists of the present invention could be used to treat or prevent blood coagulation diseases, disorders, and/or conditions (e.g., afibrinogenemia, factor deficiencies, hemophilia), blood platelet diseases, disorders, and/or conditions (e.g., thrombocytopenia), or wounds resulting from trauma, surgery, or other causes. Alternatively, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention that can decrease hemostatic or thrombolytic activity could be used to inhibit or dissolve clotting. These molecules could be important in the treatment or prevention of heart attacks (infarction), strokes, or scarring.

[0578] In specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to prevent, diagnose, prognose, and/or treat thrombosis, arterial thrombosis, venous thrombosis,

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thromboembolism, pulmonary embolism, atherosclerosis, myocardial infarction, transient ischemic attack, unstable angina. In specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used for the prevention of occulsion of saphenous grafts, for reducing the risk of periprocedural thrombosis as might accompany angioplasty procedures, for reducing the risk of stroke in patients with atrial fibrillation including nonrheumatic atrial fibrillation, for reducing the risk of embolism associated with mechanical heart valves and or mitral valves disease. Other uses for the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention, include, but are not limited to, the prevention of occlusions in extrcorporeal devices (e.g., intravascular canulas, vascular access shunts in hemodialysis patients, hemodialysis machines, and cardiopulmonary bypass machines).

[0579] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to prevent, diagnose, prognose, and/or treat diseases and disorders of the blood and/or blood forming organs associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1A, column 7 (Tissue Distribution Library Code).

[0580] polynucleotides, polypeptides, antibodies, and/or agonists or The antagonists of the present invention may be used to modulate hematopoietic activity (the formation of blood cells). For example, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to increase the quantity of all or subsets of blood cells, such as, for example, erythrocytes, lymphocytes (B or T cells), myeloid cells (e.g., basophils, eosinophils, neutrophils, mast cells, macrophages) and platelets. The ability to decrease the quantity of blood cells or subsets of blood cells may be useful in the prevention, detection, diagnosis and/or treatment of anemias and leukopenias described below. Alternatively, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to decrease the quantity of all or subsets of blood cells, such as, for example, erythrocytes, lymphocytes (B or T cells), myeloid cells (e.g., basophils, eosinophils, neutrophils, mast cells, macrophages) and platelets.. The ability to decrease the quantity of blood cells or subsets of blood cells

may be useful in the prevention, detection, diagnosis and/or treatment of leukocytoses, such as, for example eosinophilia.

[0581] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to prevent, treat, or diagnose blood dyscrasia.

[0582] Anemias are conditions in which the number of red blood cells or amount of hemoglobin (the protein that carries oxygen) in them is below normal. Anemia may be caused by excessive bleeding, decreased red blood cell production, or increased red blood cell destruction (hemolysis). The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias. Anemias that may be treated prevented or diagnosed by the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include iron deficiency anemia, hypochromic anemia, microcytic anemia, chlorosis, hereditary siderob;astic anemia, idiopathic acquired sideroblastic anemia, red cell aplasia, megaloblastic anemia (e.g., pernicious anemia, (vitamin B12 deficiency) and folic acid deficiency anemia), aplastic anemia, hemolytic anemias (e.g., autoimmune helolytic anemia, microangiopathic hemolytic anemia, and paroxysmal nocturnal hemoglobinuria). The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias associated with diseases including but not limited to, anemias associated with systemic lupus erythematosus, cancers, lymphomas, chronic renal disease, and enlarged spleens. The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias arising from drug treatments such as anemias associated with methyldopa, dapsone, and/or sulfadrugs. Additionally, rhe polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias associated with abnormal red blood cell architecture including but not limited to, hereditary spherocytosis, hereditary elliptocytosis, glucose-6-phosphate dehydrogenase deficiency, and sickle cell anemia.

[0583] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or

diagnosing hemoglobin abnormalities, (e.g., those associated with sickle cell anemia, hemoglobin C disease, hemoglobin S-C disease, and hemoglobin E disease). Additionally, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating thalassemias, including, but not limited to major and minor forms of alpha-thalassemia and beta-thalassemia.

In another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating bleeding disorders including, but not limited to, thrombocytopenia (e.g., idiopathic thrombocytopenic purpura, and thrombotic thrombocytopenic purpura), Von Willebrand's disease, hereditary platelet disorders (e.g., storage pool disease such as Chediak-Higashi and Hermansky-Pudlak syndromes, thromboxane A2 dysfunction, thromboasthenia, and Bernard-Soulier syndrome), hemolytic-uremic syndrome, hemophelias such as hemophelia A or Factor VII deficiency and Christmas disease or Factor IX deficiency, Hereditary Hemorhhagic Telangiectsia, also known as Rendu-Osler-Weber syndrome, allergic purpura (Henoch Schonlein purpura) and disseminated intravascular coagulation.

The effect of the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention on the clotting time of blood may be monitored using any of the clotting tests known in the art including, but not limited to, whole blood partial thromboplastin time (PTT), the activated partial thromboplastin time (aPTT), the activated clotting time (ACT), the recalcified activated clotting time, or the Lee-White Clotting time.

[0586] Several diseases and a variety of drugs can cause platelet dysfunction. Thus, in a specific embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating acquired platelet dysfunction such as platelet dysfunction accompanying kidney failure, leukemia, multiple myeloma, cirrhosis of the liver, and systemic lupus erythematosus as well as platelet dysfunction associated with drug treatments, including treatment with aspirin, ticlopidine, nonsteroidal anti-inflammatory drugs (used for arthritis, pain, and sprains), and penicillin in high doses.

[0587] In another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders characterized by or associated with increased or decreased numbers of white blood cells. Leukopenia occurs when the number of white blood cells decreases below normal. Leukopenias include, but are not limited to, neutropenia and lymphocytopenia. An increase in the number of white blood cells compared to normal is known as leukocytosis. The body generates increased numbers of white blood cells during infection. Thus, leukocytosis may simply be a normal physiological parameter that reflects infection. Alternatively, leukocytosis may be an indicator of injury or other disease such as cancer. Leokocytoses, include but are not limited to, eosinophilia, and accumulations of macrophages. In specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating leukopenia. In other specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating leukocytosis

[0588] Leukopenia may be a generalized decreased in all types of white blood cells, or may be a specific depletion of particular types of white blood cells. Thus, in specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating decreases in neutrophil numbers, known as neutropenia. Neutropenias that may be diagnosed, prognosed, prevented, and/or treated by the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include, but are not limited to, infantile genetic agranulocytosis, familial neutropenia, cyclic neutropenia, neutropenias resulting from or associated with dietary deficiencies (e.g., vitamin B 12 deficiency or folic acid deficiency), neutropenias resulting from or associated with drug treatments (e.g., antibiotic regimens such as penicillin treatment, sulfonamide treatment, anticoagulant treatment, anticonvulsant drugs, anti-thyroid drugs, and cancer chemotherapy), and neutropenias resulting from increased neutrophil destruction that may occur in association with some bacterial or viral infections, allergic disorders, autoimmune diseases, conditions in which an

individual has an enlarged spleen (e.g., Felty syndrome, malaria and sarcoidosis), and some drug treatment regimens.

antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating lymphocytopenias (decreased numbers of B and/or T lymphocytes), including, but not limited lymphocytopenias resulting from or associated with stress, drug treatments (e.g., drug treatment with corticosteroids, cancer chemotherapies, and/or radiation therapies), AIDS infection and/or other diseases such as, for example, cancer, rheumatoid arthritis, systemic lupus erythematosus, chronic infections, some viral infections and/or hereditary disorders (e.g., DiGeorge syndrome, Wiskott-Aldrich Syndome, severe combined immunodeficiency, ataxia telangiectsia).

[0590] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders associated with macrophage numbers and/or macrophage function including, but not limited to, Gaucher's disease, Niemann-Pick disease, Letterer-Siwe disease and Hand-Schuller-Christian disease.

[0591] In another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders associated with eosinophil numbers and/or eosinophil function including, but not limited to, idiopathic hypereosinophilic syndrome, eosinophilia-myalgia syndrome, and Hand-Schuller-Christian disease.

In yet another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating leukemias and lymphomas including, but not limited to, acute lymphocytic (lymphpblastic) leukemia (ALL), acute myeloid (myelocytic, myelogenous, myeloblastic, or myelomonocytic) leukemia, chronic lymphocytic leukemia (e.g., B cell leukemias, T cell leukemias, Sezary syndrome, and Hairy cell leukenia), chronic myelocytic (myeloid, myelogenous, or granulocytic) leukemia, Hodgkin's lymphoma, non-hodgkin's lymphoma, Burkitt's lymphoma, and mycosis fungoides.

[0593] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders of plasma cells including, but not limited to, plasma cell dyscrasias, monoclonal gammaopathies, monoclonal gammopathies of undetermined significance, multiple myeloma, macroglobulinemia, Waldenstrom's macroglobulinemia, cryoglobulinemia, and Raynaud's phenomenon.

- [0594] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing myeloproliferative disorders, including but not limited to, polycythemia vera, relative polycythemia, secondary polycythemia, myelofibrosis, acute myelofibrosis, agnogenic myelod metaplasia, thrombocythemia, (including both primary and seconday thrombocythemia) and chronic myelocytic leukemia.
- [0595] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as a treatment prior to surgery, to increase blood cell production.
- [0596] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to enhance the migration, phagocytosis, superoxide production, antibody dependent cellular cytotoxicity of neutrophils, eosionophils and macrophages.
- [0597] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to increase the number of stem cells in circulation prior to stem cells pheresis. In another specific embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to increase the number of stem cells in circulation prior to platelet pheresis.
- [0598] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to increase cytokine production.
- [0599] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in preventing, diagnosing, and/or treating primary hematopoietic disorders.

Hyperproliferative Disorders

[0600] Reproductive system associated polynucleotides or polypeptides, or agonists or antagonists thereof, can be used to treat, prevent, diagnose and/or prognose hyperproliferative diseases, disorders, and/or conditions, including neoplasms.

[0601] In a specific embodiment, reproductive system associated polynucleotides or polypeptides, or agonists or antagonists thereof, can be used to treat, prevent, and/or diagnose hyperproliferative diseases, disorders, and/or conditions of the reproductive system.

[0602] In a preferred embodiment, reproductive system associated polynucleotides or polypeptides, or agonists or antagonists thereof, can be used to treat, prevent, and/or diagnose reproductive system neoplasms.

[0603] Reproductive system associated polynucleotides or polypeptides, or agonists or antagonists of the invention, may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, reproductive system associated polynucleotides or polypeptides, or agonists or antagonists thereof, may proliferate other cells, which can inhibit the hyperproliferative disorder.

[0604] For example, by increasing an immune response, particularly increasing antigenic qualities of the hyperproliferative disorder or by proliferating, differentiating, or mobilizing T-cells, hyperproliferative diseases, disorders, and/or conditions can be treated, prevented, and/or diagnosed. This immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, decreasing an immune response may also be a method of treating, preventing, and/or diagnosing hyperproliferative diseases, disorders, and/or conditions, such as a chemotherapeutic agent.

[0605] Examples of hyperproliferative diseases, disorders, and/or conditions that can be treated, prevented, and/or diagnosed by reproductive system associated polynucleotides or polypeptides, or agonists or antagonists thereof, include, but are not limited to neoplasms located in the: prostate, colon, abdomen, bone, breast, digestive system, liver, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous (central and peripheral), lymphatic system, pelvic, skin, soft tissue, spleen, thoracic, and urogenital.

[0606] Similarly, other hyperproliferative disorders can also be treated or detected by polynucleotides or polypeptides, or agonists or antagonists of the present invention. Examples of such hyperproliferative disorders include, but are not limited to: Acute Childhood Lymphoblastic Leukemia, Acute Lymphoblastic Leukemia, Acute Lymphocytic Leukemia, Acute Myeloid Leukemia, Adrenocortical Carcinoma, Adult (Primary) Hepatocellular Cancer, Adult (Primary) Liver Cancer, Adult Acute Lymphocytic Leukemia, Adult Acute Myeloid Leukemia, Adult Hodgkin's Disease, Adult Hodgkin's Lymphoma, Adult Lymphocytic Leukemia, Adult Non-Hodgkin's Lymphoma, Adult Primary Liver Cancer, Adult Soft Tissue Sarcoma, AIDS-Related Lymphoma, AIDS-Related Malignancies, Anal Cancer, Astrocytoma, Bile Duct Cancer, Bladder Cancer, Bone Cancer, Brain Stem Glioma, Brain Tumors, Breast Cancer, Cancer of the Renal Pelvis and Ureter, Central Nervous System (Primary) Lymphoma, Central Nervous System Lymphoma, Cerebellar Astrocytoma, Cerebral Astrocytoma, Cervical Cancer, Childhood (Primary) Hepatocellular Cancer, Childhood (Primary) Liver Cancer, Childhood Acute Lymphoblastic Leukemia, Childhood Acute Myeloid Leukemia, Childhood Brain, Stem Glioma, Childhood Cerebellar Astrocytoma, Childhood Cerebral Astrocytoma, Childhood Extracranial Germ Cell Tumors, Childhood Hodgkin's Disease, Childhood Hodgkin's Lymphoma, Childhood Hypothalamic and Visual Pathway Glioma, Childhood Lymphoblastic Leukemia, Childhood Medulloblastoma, Childhood Non-Hodgkin's Lymphoma, Childhood Pineal and Supratentorial Primitive Neuroectodermal Tumors, Childhood Primary Liver Cancer, Childhood Rhabdomyosarcoma, Childhood Soft Tissue Childhood Visual Pathway and Hypothalamic Glioma, Chronic Sarcoma, Lymphocytic Leukemia, Chronic Myelogenous Leukemia, Colon Cancer, Cutaneous T-Cell Lymphoma, Endocrine Pancreas Islet Cell Carcinoma, Endometrial Cancer, Ependymoma, Epithelial Cancer, Esophageal Cancer, Ewing's Sarcoma and Related Tumors, Exocrine Pancreatic Cancer, Extracranial Germ Cell Tumor, Extragonadal Germ Cell Tumor, Extrahepatic Bile Duct Cancer, Eye Cancer, Female Breast Cancer, Gaucher's Disease, Gallbladder Cancer, Gastric Cancer, Gastrointestinal Carcinoid Tumor, Gastrointestinal Tumors, Germ Cell Tumors, Gestational Trophoblastic Tumor, Hairy Cell Leukemia, Head and Neck Cancer, Hepatocellular Cancer, Hodgkin's Disease, Hodgkin's Lymphoma. Hypergammaglobulinemia,

Hypopharyngeal Cancer, Intestinal Cancers, Intraocular Melanoma, Islet Cell Carcinoma, Islet Cell Pancreatic Cancer, Kaposi's Sarcoma, Kidney Cancer, Laryngeal Cancer, Lip and Oral Cavity Cancer, Liver Cancer, Lung Cancer, Lymphoproliferative Disorders, Macroglobulinemia, Male Breast Cancer, Malignant Mesothelioma, Malignant Thymoma, Medulloblastoma, Melanoma, Mesothelioma, Metastatic Occult Primary Squamous Neck Cancer, Metastatic Primary Squamous Neck Cancer, Metastatic Squamous Neck Cancer, Multiple Myeloma, Multiple Myeloma/Plasma Cell Neoplasm, Myelodysplastic Syndrome, Myelogenous Leukemia, Myeloid Leukemia, Myeloproliferative Disorders, Nasal Cavity and Paranasal Sinus Cancer, Nasopharyngeal Cancer, Neuroblastoma, Non-Hodgkin's Lymphoma During Pregnancy, Nonmelanoma Skin Cancer, Non-Small Cell Lung Cancer, Occult Primary Metastatic Squamous Neck Cancer, Oropharyngeal Cancer, Osteo-/Malignant Fibrous Sarcoma, Osteosarcoma/Malignant Fibrous Histiocytoma, Osteosarcoma/Malignant Fibrous Histiocytoma of Bone, Ovarian Epithelial Cancer, Ovarian Germ Cell Tumor, Ovarian Low Malignant Potential Tumor, Pancreatic Cancer, Paraproteinemias, Purpura, Parathyroid Cancer, Penile Cancer, Pheochromocytoma, Pituitary Tumor, Plasma Cell Neoplasm/Multiple Myeloma, Primary Central Nervous System Lymphoma, Primary Liver Cancer, Prostate Cancer, Rectal Cancer, Renal Cell Cancer, Renal Pelvis and Ureter Cancer, Retinoblastoma, Rhabdomyosarcoma, Salivary Gland Cancer, Sarcoidosis Sarcomas, Sezary Syndrome, Skin Cancer, Small Cell Lung Cancer, Small Intestine Cancer, Soft Tissue Sarcoma, Squamous Neck Cancer, Stomach Cancer, Supratentorial Primitive Neuroectodermal and Pineal Tumors, T-Cell Lymphoma, Testicular Cancer, Thymoma, Thyroid Cancer, Transitional Cell Cancer of the Renal Pelvis and Ureter, Transitional Renal Pelvis and Ureter Cancer, Trophoblastic Tumors, Ureter and Renal Pelvis Cell Cancer, Urethral Cancer, Uterine Cancer, Uterine Sarcoma, Vaginal Cancer, Visual Pathway and Hypothalamic Glioma, Vulvar Cancer, Waldenstrom's Macroglobulinemia, Wilms' Tumor, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.

[0607] In another preferred embodiment, polynucleotides or polypeptides, or agonists or antagonists of the present invention are used to diagnose, prognose, prevent, and/or treat premalignant conditions and to prevent progression to a neoplastic or malignant state, including but not limited to those disorders described

above. Such uses are indicated in conditions known or suspected of preceding progression to neoplasia or cancer, in particular, where non-neoplastic cell growth consisting of hyperplasia, metaplasia, or most particularly, dysplasia has occurred (for review of such abnormal growth conditions, see Robbins and Angell, 1976, Basic Pathology, 2d Ed., W. B. Saunders Co., Philadelphia, pp. 68-79.)

[0608] Hyperplasia is a form of controlled cell proliferation, involving an increase in cell number in a tissue or organ, without significant alteration in structure or function. Hyperplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, angiofollicular mediastinal lymph node hyperplasia, angiolymphoid hyperplasia with eosinophilia, atypical melanocytic hyperplasia, basal cell hyperplasia, benign giant lymph node hyperplasia, cementum hyperplasia, congenital adrenal hyperplasia, congenital sebaceous hyperplasia, cystic hyperplasia, cystic hyperplasia of the breast, denture hyperplasia, ductal hyperplasia, endometrial hyperplasia, fibromuscular hyperplasia, focal epithelial hyperplasia, gingival hyperplasia, inflammatory fibrous hyperplasia, inflammatory papillary hyperplasia, intravascular papillary endothelial hyperplasia, nodular hyperplasia prostate. nodular regenerative hyperplasia, pseudoepitheliomatous hyperplasia, senile sebaceous hyperplasia, and verrucous hyperplasia.

[0609] Metaplasia is a form of controlled cell growth in which one type of adult or fully differentiated cell substitutes for another type of adult cell. Metaplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, agnogenic myeloid metaplasia, apocrine metaplasia, atypical metaplasia, autoparenchymatous metaplasia, connective tissue metaplasia, epithelial metaplasia, intestinal metaplasia, metaplastic anemia, metaplastic ossification, metaplastic polyps, myeloid metaplasia, primary myeloid metaplasia, secondary myeloid metaplasia, squamous metaplasia, squamous metaplasia of amnion, and symptomatic myeloid metaplasia.

[0610] Dysplasia is frequently a forerunner of cancer, and is found mainly in the epithelia; it is the most disorderly form of non-neoplastic cell growth, involving a loss

in individual cell uniformity and in the architectural orientation of cells. Dysplastic cells often have abnormally large, deeply stained nuclei, and exhibit pleomorphism. Dysplasia characteristically occurs where there exists chronic irritation or inflammation. Dysplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, anhidrotic ectodermal dysplasia, anterofacial dysplasia, asphyxiating thoracic dysplasia, atriodigital dysplasia, bronchopulmonary dysplasia, cerebral dysplasia, cervical dysplasia, chondroectodermal dysplasia, cleidocranial dysplasia, congenital ectodermal dysplasia, craniodiaphysial dysplasia, craniocarpotarsal dysplasia, craniometaphysial dysplasia, dentin dysplasia, diaphysial dysplasia, ectodermal dysplasia, enamel dysplasia, encephalo-ophthalmic dysplasia, dysplasia epiphysialis hemimelia, dysplasia epiphysialis multiplex, dysplasia epiphysialis punctata, epithelial dysplasia, faciodigitogenital dysplasia, familial fibrous dysplasia of jaws, familial white folded dysplasia, fibromuscular dysplasia, fibrous dysplasia of bone, florid osseous dysplasia, hereditary renal-retinal dysplasia, hidrotic ectodermal dysplasia, hypohidrotic ectodermal dysplasia, lymphopenic thymic dysplasia, mammary dysplasia, mandibulofacial dysplasia, metaphysial dysplasia, Mondini dysplasia, monostotic fibrous dysplasia, mucoepithelial dysplasia, multiple epiphysial dysplasia, oculoauriculovertebral dysplasia, oculodentodigital dysplasia, oculovertebral dysplasia, odontogenic dysplasia, ophthalmomandibulomelic dysplasia, periapical cemental dysplasia, polyostotic fibrous dysplasia, pseudoachondroplastic spondyloepiphysial dysplasia, retinal dysplasia, septo-optic dysplasia, spondyloepiphysial dysplasia, and ventriculoradial dysplasia.

[0611] Additional pre-neoplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, benign dysproliferative disorders (e.g., benign tumors, fibrocystic conditions, tissue hypertrophy, intestinal polyps, colon polyps, and esophageal dysplasia), leukoplakia, keratoses, Bowen's disease, Farmer's Skin, solar cheilitis, and solar keratosis.

[0612] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to

diagnose and/or prognose disorders associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1A, 7 (Tissue Distribution Library Code).

In another embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention conjugated to a toxin or a radioactive isotope, as described herein, may be used to treat cancers and neoplasms, including, but not limited to those described herein. In a further preferred embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention conjugated to a toxin or a radioactive isotope, as described herein, may be used to treat acute myelogenous leukemia.

Additionally, polynucleotides, polypeptides, and/or agonists or antagonists of the invention may affect apoptosis, and therefore, would be useful in treating a number of diseases associated with increased cell survival or the inhibition of apoptosis. For example, diseases associated with increased cell survival or the inhibition of apoptosis that could be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include cancers (such as follicular lymphomas, carcinomas with p53 mutations, and hormone-dependent tumors, including, but not limited to colon cancer, cardiac tumors, pancreatic cancer, melanoma, retinoblastoma, glioblastoma, lung cancer, intestinal cancer, testicular cancer, stomach cancer, neuroblastoma, myxoma, myoma, lymphoma. endothelioma, osteoblastoma, osteoclastoma, osteosarcoma, chondrosarcoma, adenoma, breast cancer, prostate cancer, Kaposi's sarcoma and ovarian cancer); autoimmune disorders such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis) and viral infections (such as herpes viruses, pox viruses and adenoviruses), inflammation, graft v. host disease, acute graft rejection, and chronic graft rejection.

[0615] In preferred embodiments, polynucleotides, polypeptides, and/or agonists or antagonists of the invention are used to inhibit growth, progression, and/or metastasis of cancers, in particular those listed above.

Additional diseases or conditions associated with increased cell survival [0616] that could be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include, but are not limited to, progression, and/or metastases of malignancies and related disorders such as leukemia (including acute leukemias (e.g., acute lymphocytic leukemia, acute myelocytic leukemia (including myeloblastic, promyelocytic, myelomonocytic, monocytic, and erythroleukemia)) and chronic leukemias (e.g., chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia)), polycythemia vera, lymphomas (e.g., Hodgkin's disease and non-Hodgkin's disease), multiple myeloma, Waldenstrom's macroglobulinemia, heavy chain disease, and solid tumors including, but not limited to, sarcomas and carcinomas such as fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, chordoma, angiosarcoma, endotheliosarcoma, lymphangiosarcoma, lymphangioendotheliosarcoma, synovioma, mesothelioma, Ewing's tumor, leiomyosarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary adenocarcinomas, cystadenocarcinoma, medullary carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma, choriocarcinoma, seminoma, embryonal carcinoma, Wilm's tumor, cervical cancer, testicular tumor, lung carcinoma, small cell lung carcinoma, bladder carcinoma, epithelial carcinoma, glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, pinealoma, emangioblastoma, acoustic neuroma, oligodendroglioma, menangioma, melanoma, neuroblastoma, and retinoblastoma.

prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include AIDS; neurodegenerative disorders (such as Alzheimer's disease, Parkinson's disease, amyotrophic lateral sclerosis, retinitis pigmentosa, cerebellar degeneration and brain tumor or prior associated disease); autoimmune disorders (such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis)

myelodysplastic syndromes (such as aplastic anemia), graft v. host disease, ischemic injury (such as that caused by myocardial infarction, stroke and reperfusion injury), liver injury (e.g., hepatitis related liver injury, ischemia/reperfusion injury, cholestosis (bile duct injury) and liver cancer); toxin-induced liver disease (such as that caused by alcohol), septic shock, cachexia and anorexia.

- [0618] Hyperproliferative diseases and/or disorders that could be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include, but are not limited to, neoplasms located in the liver, abdomen, bone, breast, digestive system, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous system (central and peripheral), lymphatic system, pelvis, skin, soft tissue, spleen, thorax, and urogenital tract.
- [0619] Similarly, other hyperproliferative disorders can also be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention. Examples of such hyperproliferative disorders include, but are not limited to: hypergammaglobulinemia, lymphoproliferative disorders, paraproteinemias, purpura, sarcoidosis, Sezary Syndrome, Waldenstron's macroglobulinemia, Gaucher's Disease, histiocytosis, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.
- [0620] One preferred embodiment utilizes polynucleotides of the present invention to inhibit aberrant cellular division, by gene therapy using the present invention, and/or protein fusions or fragments thereof.
- [0621] Thus, the present invention provides a method for treating cell proliferative diseases, disorders, and/or conditions by inserting into an abnormally proliferating cell a polynucleotide of the present invention, wherein said polynucleotide represses said cell proliferation, disease, disorder, and/or condition.
- [0622] In a preferred embodiment, the present invention provides a method for treating cell proliferative diseases, disorders and/or conditions of the reproductive system by inserting into a cell, a polynucleotide of the present invention, wherein said polynucleotide represses said cell proliferation, disease and/or disorder.
- [0623] Another embodiment of the present invention provides a method of treating cell-proliferative diseases, disorders, and/or conditions in individuals comprising

administration of one or more active gene copies of the present invention to an abnormally proliferating cell or cells. In a preferred embodiment, polynucleotides of the present invention is a DNA construct comprising a recombinant expression vector effective in expressing a DNA sequence encoding said polynucleotides. In another preferred embodiment of the present invention, the DNA construct encoding the polynucleotides of the present invention is inserted into cells to be treated utilizing a retrovirus, or more preferably an adenoviral vector (see, e.g., G J. Nabel, et. al., PNAS 96: 324-326 (1999), which is hereby incorporated by reference). In a most preferred embodiment, the viral vector is defective and will not transform non-proliferating cells, only proliferating cells. Moreover, in a preferred embodiment, the polynucleotides of the present invention inserted into proliferating cells either alone, or in combination with or fused to other polynucleotides, can then be modulated via an external stimulus (i.e., magnetic, specific small molecule, chemical, or drug administration, etc.), which acts upon the promoter upstream of said polynucleotides to induce expression of the encoded protein product. As such the beneficial therapeutic affect of the present invention may be expressly modulated (i.e., to increase, decrease, or inhibit expression of the present invention) based upon said external stimulus.

[0624] Polynucleotides of the present invention may be useful in repressing expression of oncogenic genes or antigens. By "repressing expression of the oncogenic genes" is intended the suppression of the transcription of the gene, the degradation of the gene transcript (pre-message RNA), the inhibition of splicing, the destruction of the messenger RNA, the prevention of the post-translational modifications of the protein, the destruction of the protein, or the inhibition of the normal function of the protein.

[0625] For local administration to abnormally proliferating cells, polynucleotides of the present invention may be administered by any method known to those of skill in the art including, but not limited to transfection, electroporation, microinjection of cells, or in vehicles such as liposomes, lipofectin, or as naked polynucleotides, or any other method described throughout the specification. The polynucleotide of the present invention may be delivered by known gene delivery systems such as, but not limited to, retroviral vectors (Gilboa, J. Virology 44:845 (1982); Hocke, Nature

320:275 (1986); Wilson, et al., Proc. Natl. Acad. Sci. U.S.A. 85:3014), vaccinia virus system (Chakrabarty et al., Mol. Cell Biol. 5:3403 (1985) or other efficient DNA delivery systems (Yates et al., Nature 313:812 (1985)) known to those skilled in the art. These references are exemplary only and are hereby incorporated by reference. In order to specifically deliver or transfect cells which are abnormally proliferating and spare non-dividing cells, it is preferable to utilize a retrovirus, or adenoviral (as described in the art and elsewhere herein) delivery system known to those of skill in the art. Since host DNA replication is required for retroviral DNA to integrate and the retrovirus will be unable to self replicate due to the lack of the retrovirus genes needed for its life cycle. Utilizing such a retroviral delivery system for polynucleotides of the present invention will target said gene and constructs to abnormally proliferating cells and will spare the non-dividing normal cells.

[0626] The polynucleotides of the present invention may be delivered directly to cell proliferative disorder/disease sites in internal organs, body cavities and the like by use of imaging devices used to guide an injecting needle directly to the disease site. The polynucleotides of the present invention may also be administered to disease sites at the time of surgical intervention.

[0627] By "cell proliferative disease" is meant any human or animal disease or disorder, affecting any one or any combination of organs, cavities, or body parts, which is characterized by single or multiple local abnormal proliferations of cells, groups of cells, or tissues, whether benign or malignant.

Any amount of the polynucleotides of the present invention may be administered as long as it has a biologically inhibiting effect on the proliferation of the treated cells. Moreover, it is possible to administer more than one of the polynucleotide of the present invention simultaneously to the same site. By "biologically inhibiting" is meant partial or total growth inhibition as well as decreases in the rate of proliferation or growth of the cells. The biologically inhibitory dose may be determined by assessing the effects of the polynucleotides of the present invention on target malignant or abnormally proliferating cell growth in tissue culture, tumor growth in animals and cell cultures, or any other method known to one of ordinary skill in the art.

[0629] The present invention is further directed to antibody-based therapies which involve administering of anti-polypeptides and anti-polynucleotide antibodies to a mammalian, preferably human, patient for treating one or more of the described diseases, disorders, and/or conditions. Methods for producing anti-polypeptides and anti-polynucleotide antibodies polyclonal and monoclonal antibodies are described in detail elsewhere herein. Such antibodies may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

[0630] A summary of the ways in which the antibodies of the present invention may be used therapeutically includes binding polynucleotides or polypeptides of the present invention locally or systemically in the body or by direct cytotoxicity of the antibody, e.g., as mediated by complement (CDC) or by effector cells (ADCC). Some of these approaches are described in more detail below. Armed with the teachings provided herein, one of ordinary skill in the art will know how to use the antibodies of the present invention for diagnostic, monitoring or therapeutic purposes without undue experimentation.

[0631] In particular, the antibodies, fragments and derivatives of the present invention are useful for treating a subject having or developing cell proliferative and/or differentiation diseases, disorders, and/or conditions as described herein. Such treatment comprises administering a single or multiple doses of the antibody, or a fragment, derivative, or a conjugate thereof.

[0632] The antibodies of this invention may be advantageously utilized in combination with other monoclonal or chimeric antibodies, or with lymphokines or hematopoietic growth factors, for example, which serve to increase the number or activity of effector cells which interact with the antibodies.

[0633] It is preferred to use high affinity and/or potent *in vivo* inhibiting and/or neutralizing antibodies against polypeptides or polynucleotides of the present invention, fragments or regions thereof, for both immunoassays directed to and therapy of diseases, disorders, and/or conditions related to polynucleotides or polypeptides, including fragments thereof, of the present invention. Such antibodies, fragments, or regions, will preferably have an affinity for polynucleotides or polypeptides, including fragments thereof. Preferred binding affinities include those with a dissociation constant or Kd less than 5X10-6M, 10-6M, 5X10-7M, 10-7M, 5X10-7M,

 8 M, 10^{-8} M, $5X10^{-9}$ M, 10^{-9} M, $5X10^{-10}$ M, 10^{-10} M, $5X10^{-11}$ M, 10^{-11} M, $5X10^{-12}$ M, 10^{-12} M, $5X10^{-13}$ M, 10^{-13} M, $5X10^{-14}$ M, 10^{-14} M, $5X10^{-15}$ M, and 10^{-15} M.

[0634] Moreover, reproductive system antigen polypeptides of the present invention or fragments thereof, are useful in inhibiting the angiogenesis of proliferative cells or tissues, either alone, as a protein fusion, or in combination with other polypeptides directly or indirectly, as described elsewhere herein. In a most preferred embodiment, said anti-angiogenesis effect may be achieved indirectly, for example, through the inhibition of hematopoietic, tumor-specific cells, such as tumor-associated macrophages (see, e.g., Joseph IB, et al. J Natl Cancer Inst, 90(21):1648-53 (1998), which is hereby incorporated by reference). Antibodies directed to polypeptides or polynucleotides of the present invention may also result in inhibition of angiogenesis directly, or indirectly (see, e.g., Witte L, et al., Cancer Metastasis Rev. 17(2):155-61 (1998), which is hereby incorporated by reference)).

Polypeptides, including protein fusions, of the present invention, or [0635] fragments thereof may be useful in inhibiting proliferative cells or tissues through the induction of apoptosis. Said polypeptides may act either directly, or indirectly to induce apoptosis of proliferative cells and tissues, for example in the activation of a death-domain receptor, such as tumor necrosis factor (TNF) receptor-1, CD95 (Fas/APO-1), TNF-receptor-related apoptosis-mediated protein (TRAMP) and TNFrelated apoptosis-inducing ligand (TRAIL) receptor-1 and -2 (see, e.g., Schulze-Osthoff K, et.al., Eur J Biochem 254(3):439-59 (1998), which is hereby incorporated by reference). Moreover, in another preferred embodiment of the present invention, said polypeptides may induce apoptosis through other mechanisms, such as in the activation of other proteins which will activate apoptosis, or through stimulating the expression of said proteins, either alone or in combination with small molecule drugs or adjuvants, such as apoptonin, galectins, thioredoxins, antiinflammatory proteins (See for example, Mutat. Res. 400(1-2):447-55 (1998), Med Hypotheses.50(5):423-33 (1998), Chem. Biol. Interact. Apr 24;111-112:23-34 (1998), J. Mo. Med. 76(6):402-12 (1998), Int. J. Tissue React. 20(1):3-15 (1998), which are all hereby incorporated by reference).

[0636] Polypeptides, including protein fusions to, or fragments thereof, of the present invention are useful in inhibiting the metastasis of proliferative cells or tissues.

Inhibition may occur as a direct result of administering polypeptides, or antibodies directed to said polypeptides as described elsewhere herein, or indirectly, such as activating the expression of proteins known to inhibit metastasis, for example alpha 4 integrins, (See, e.g., Curr Top Microbiol Immunol 1998;231:125-41, which is hereby incorporated by reference). Such therapeutic affects of the present invention may be achieved either alone, or in combination with small molecule drugs or adjuvants.

[0637] In another embodiment, the invention provides a method of delivering compositions containing the polypeptides of the invention (e.g., compositions containing polypeptides or anti-reproductive system antigen polypeptide antibodies associated with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs) to targeted cells expressing the polypeptide of the present invention: reproductive system antigen polypeptides or anti-reproductive system antigen polypeptide antibodies of the invention may be associated with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs via hydrophobic, hydrophilic, ionic and/or covalent interactions.

[0638] Polypeptides, protein fusions to, or fragments thereof, of the present invention are useful in enhancing the immunogenicity and/or antigenicity of proliferating cells or tissues, either directly, such as would occur if the polypeptides of the present invention 'vaccinated' the immune response to respond to proliferative antigens and immunogens, or indirectly, such as in activating the expression of proteins known to enhance the immune response (e.g. chemokines), to said antigens and immunogens.

Urinary System Disorders

[0639] Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose disorders of the urinary system, including but not limited to disorders of the renal system, bladder, ureters, and urethra. Renal disorders include, but are not limited to, kidney failure, nephritis, blood vessel disorders of kidney, metabolic and congenital kidney disorders, urinary disorders of the kidney, autoimmune disorders, sclerosis and necrosis, electrolyte imbalance, and kidney cancers.

[0640] Kidney failure diseases include, but are not limited to, acute kidney failure, chronic kidney failure, atheroembolic renal failure, and end-stage renal disease. Inflammatory diseases of the kidney include acute glomerulonephritis, postinfectious glomerulonephritis, rapidly progressive glomerulonephritis, nephrotic syndrome, membranous glomerulonephritis, familial nephrotic syndrome, membranoproliferative glomerulonephritis I and II, mesangial proliferative glomerulonephritis, chronic glomerulonephritis, acute tubulointerstitial nephritis, chronic tubulointerstitial nephritis, acute post-streptococcal glomerulonephritis (PSGN), pyelonephritis, lupus nephritis, chronic nephritis, interstitial nephritis, and post-streptococcal glomerulonephritis.

[0641] Blood vessel disorders of the kidneys include, but are not limited to, kidney infarction, atheroembolic kidney disease, cortical necrosis, malignant nephrosclerosis, renal vein thrombosis, renal underperfusion, renal ischemia-reperfusion, renal artery embolism, and renal artery stenosis. Kidney disorders resulting form urinary tract problems include, but are not limited to, pyelonephritis, hydronephrosis, urolithiasis (renal lithiasis, nephrolithiasis), reflux nephropathy, urinary tract infections, urinary retention, and acute or chronic unilateral obstructive uropathy.

limited to, renal tubular acidosis, renal glycosuria, nephrogenic diabetes insipidus, cystinuria, Fanconi's syndrome, vitamin D-resistant rickets, Hartnup disease, Bartter's syndrome, Liddle's syndrome, polycystic kidney disease, medullary cystic disease, medullary sponge kidney, Alport's syndrome, nail-patella syndrome, congenital nephrotic syndrome, CRUSH syndrome, horseshoe kidney, diabetic nephropathy, nephrogenic diabetes insipidus, analgesic nephropathy, kidney stones, and membranous nephropathy, Kidney disorders resulting from an autoimmune response include, but are not limited to, systemic lupus erythematosus (SLE), Goodpasture syndrome, IgA nephropathy, and IgM mesangial proliferative glomerulonephritis.

[0643] Sclerotic or necrotic disorders of the kidney include, but are not limited to, glomerulosclerosis, diabetic nephropathy, focal segmental glomerulosclerosis (FSGS), necrotizing glomerulonephritis, and renal papillary necrosis. Kidneys may also develop carcinomas, including, but not limited to, hypernephroma, nephroblastoma, renal cell cancer, transitional cell cancer, squamous cell cancer, and Wilm's tumor.

[0644] Kidney disorders may also result in electrolyte imbalances, including, but not limited to, nephrocalcinosis, pyuria, edema, hydronephritis, proteinuria, hyponatremia, hypernatremia, hypokalemia, hyperkalemia, hypercalcemia, hypophosphatemia, and hyperphosphatemia.

[0645] Bladder disorders include, but are not limited to, benign prostatic hyperplasia (BPH), interstitial cystitis (IC), prostatitis, proteinuria, urinary tract infections, urinary incontinence, urinary retention. Disorders of the ureters and urethra include, but are not limited to, acute or chronic unilateral obstructive uropathy. The bladder, ureters, and urethra may also develop carcinomas, including, but not limited to, superficial bladder cancer, invasive bladder cancer, carcinoma of the ureter, and urethra cancers.

[0646] Polypeptides may be administered using any method known in the art, including, but not limited to, direct needle injection at the delivery site, intravenous injection, topical administration, catheter infusion, biolistic injectors, particle accelerators, gelfoam sponge depots, other commercially available depot materials, osmotic pumps, oral or suppositorial solid pharmaceutical formulations, decanting or topical applications during surgery, aerosol delivery. Such methods are known in the art. Polypeptides may be administered as part of a Therapeutic, described in more detail below. Methods of delivering polynucleotides are described in more detail herein.

Cardiovascular Disorders

[0647] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose cardiovascular disorders, including, but not limited to, peripheral artery disease, such as limb ischemia.

[0648] Cardiovascular disorders include cardiovascular abnormalities, such as arterio-arterial fistula, arteriovenous fistula, cerebral arteriovenous malformations, congenital heart defects, pulmonary atresia, and Scimitar Syndrome. Congenital heart defects include aortic coarctation, cor triatriatum, coronary vessel anomalies, crisscross heart, dextrocardia, patent ductus arteriosus, Ebstein's anomaly, Eisenmenger complex, hypoplastic left heart syndrome, levocardia, tetralogy of fallot,

transposition of great vessels, double outlet right ventricle, tricuspid atresia, persistent truncus arteriosus, total anomalous pulmonary venous connection, hypoplastic left heart syndrome, and heart septal defects, such as aortopulmonary septal defect, endocardial cushion defects, Lutembacher's Syndrome, atrioventricular canal defect, trilogy of Fallot, ventricular heart septal defects.

Cardiovascular disorders also include heart disease, such as arrhythmias, [0649] carcinoid heart disease, high cardiac output, low cardiac output, cardiac tamponade, endocarditis (including bacterial), heart aneurysm, cardiac arrest, sudden cardiac death, congestive heart failure, congestive cardiomyopathy, paroxysmal dyspnea, cardiac edema, heart hypertrophy, congestive cardiomyopathy, left ventricular hypertrophy, right ventricular hypertrophy, post-infarction heart rupture, ventricular septal rupture, heart valve diseases, myocardial diseases, myocardial ischemia, pericardial pericarditis (including effusion. constrictive and tuberculous), pneumopericardium, postpericardiotomy syndrome, pulmonary heart disease, rheumatic heart disease, ventricular dysfunction, hyperemia, cardiovascular pregnancy complications, Scimitar Syndrome, diastolic dysfunction, enlarged heart, heart block, J-curve phenomenon, rheumatic heart disease, Marfan syndrome, cardiovascular syphilis, and cardiovascular tuberculosis.

[0650] Arrhythmias include sinus arrhythmia, atrial fibrillation, atrial flutter, bradycardia, extrasystole, Adams-Stokes Syndrome, bundle-branch block, sinoatrial block, long QT syndrome, parasystole, Lown-Ganong-Levine Syndrome, Mahaimtype pre-excitation syndrome, Wolff-Parkinson-White syndrome, sick sinus syndrome, tachycardias, and ventricular fibrillation. Tachycardias include paroxysmal tachycardia, supraventricular tachycardia, accelerated idioventricular rhythm, atrioventricular nodal reentry tachycardia, ectopic atrial tachycardia, ectopic junctional tachycardia, sinoatrial nodal reentry tachycardia, sinus tachycardia, Torsades de Pointes, and ventricular tachycardia.

[0651] Heart valve disease include aortic valve insufficiency, aortic valve stenosis, heart murmurs, aortic valve prolapse, mitral valve prolapse, tricuspid valve prolapse, mitral valve insufficiency, mitral valve stenosis, pulmonary atresia, pulmonary valve insufficiency, pulmonary valve stenosis, tricuspid atresia, tricuspid valve insufficiency, tricuspid valve stenosis, and bicuspid aortic valve.

[0652] Myocardial diseases include alcoholic cardiomyopathy, congestive cardiomyopathy, hypertrophic cardiomyopathy, aortic subvalvular stenosis, pulmonary subvalvular stenosis, restrictive cardiomyopathy, Chagas cardiomyopathy, endocardial fibroelastosis, endomyocardial fibrosis, Kearns Syndrome, Barth syndrome, myocardial reperfusion injury, and myocarditis.

- [0653] Myocardial ischemias include coronary disease, such as angina pectoris, Prinzmetal's angina, unstable angina, coronary aneurysm, coronary arteriosclerosis, coronary thrombosis, coronary vasospasm, myocardial infarction and myocardial stunning.
- Cardiovascular diseases also include vascular diseases such as aneurysms, [0654] angiodysplasia, angiomatosis, bacillary angiomatosis, Hippel-Lindau Disease, Klippel-Trenaunay-Weber Syndrome, Sturge-Weber Syndrome, angioneurotic edema, aortic diseases, Takayasu's Arteritis, aortitis, Leriche's Syndrome, arterial occlusive diseases, arteritis, enarteritis, polyarteritis nodosa, cerebrovascular disorders, diabetic angiopathies, diabetic retinopathy, embolisms, thrombosis. erythromelalgia, hemorrhoids, hepatic veno-occlusive disease, hypertension, hypotension (shock), ischemia, peripheral vascular diseases, phlebitis, superficial phlebitis, pulmonary veno-occlusive disease, chronic obstructive pulmonary disease, Buerger's disease, Raynaud's disease, CREST syndrome, retinal vein occlusion, Scimitar syndrome, superior vena cava syndrome, telangiectasia, atacia telangiectasia, hereditary hemorrhagic telangiectasia, deep vein thrombosis, varicocele, varicose veins, varicose ulcer, vasculitis, and venous insufficiency.
- [0655] Aneurysms include dissecting aneurysms, false aneurysms, infected aneurysms, ruptured aneurysms, aortic aneurysms, cerebral aneurysms, coronary aneurysms, heart aneurysms, and iliac aneurysms.
- [0656] Arterial occlusive diseases include arteriosclerosis, arteriolosclerosis, atherosclerosis, intermittent claudication, carotid stenosis, fibromuscular dysplasias, mesenteric vascular occlusion, Moyamoya disease, renal artery obstruction, retinal artery occlusion, and thromboangiitis obliterans.
- [0657] Cerebrovascular disorders include carotid artery diseases, cerebral amyloid angiopathy, cerebral aneurysm, cerebral anoxia, cerebral arteriosclerosis, cerebral arteriovenous malformation, cerebral artery diseases, cerebral embolism and

thrombosis, carotid artery thrombosis, sinus thrombosis, Wallenberg's syndrome, cerebral hemorrhage, epidural hematoma, subdural hematoma, subaraxhnoid hemorrhage, cerebral infarction, cerebral ischemia (including transient), subclavian steal syndrome, periventricular leukomalacia, vascular headache, cluster headache, migraine, and vertebrobasilar insufficiency.

[0658] Embolisms include air embolisms, amniotic fluid embolisms, cholesterol embolisms, blue toe syndrome, fat embolisms, pulmonary embolisms, and thromoboembolisms. Thrombosis include coronary thrombosis, hepatic vein thrombosis, deep vein thrombosis, retinal vein occlusion, carotid artery thrombosis, sinus thrombosis, Wallenberg's syndrome, and thrombophlebitis.

[0659] Ischemia includes cerebral ischemia, ischemic colitis, silent ischemia, compartment syndromes, anterior compartment syndrome, myocardial ischemia, reperfusion injuries, and peripheral limb ischemia. Vasculitis includes aortitis, arteritis, Behcet's Syndrome, Churg-Strauss Syndrome, mucocutaneous lymph node syndrome, thromboangiitis obliterans, hypersensitivity vasculitis, Schoenlein-Henoch purpura, allergic cutaneous vasculitis, and Wegener's granulomatosis.

[0660] Cardiovascular diseases can also occur due to electrolyte imbalances that include, but are not limited to hyponatremia, hypernatremia, hyporalemia, hyperkalemia, hypocaleemia, hypercaleemia, hypophosphatemia, and hyperphophatemia. Neoplasm and/or cancers of the cardiovascular system include, but are not limited to, myxomas, fibromas, and rhabdomyomas.

[0661] Polypeptides may be administered using any method known in the art, including, but not limited to, direct needle injection at the delivery site, intravenous injection, topical administration, catheter infusion, biolistic injectors, particle accelerators, gelfoam sponge depots, other commercially available depot materials, osmotic pumps, oral or suppositorial solid pharmaceutical formulations, decanting or topical applications during surgery, aerosol delivery. Such methods are known in the art. Polypeptides may be administered as part of a Therapeutic, described in more detail below. Methods of delivering polynucleotides are described in more detail herein.

Respiratory Disorders

[0662] Polynucleotides or polypeptides, or agonists or antagonists of the present invention may be used to treat, prevent, diagnose, and/or prognose diseases and/or disorders of the respiratory system.

Diseases and disorders of the respiratory system include, but are not limited 106631 to, nasal vestibulitis, nonallergic rhinitis (e.g., acute rhinitis, chronic rhinitis, atrophic rhinitis, vasomotor rhinitis), nasal polyps, and sinusitis, juvenile angiofibromas, cancer of the nose and juvenile papillomas, vocal cord polyps, nodules (singer's nodules), contact ulcers, vocal cord paralysis, laryngoceles, pharyngitis (e.g., viral and bacterial), tonsillitis, tonsillar cellulitis, parapharyngeal abscess, laryngoceles, and throat cancers (e.g., cancer of the nasopharynx, tonsil cancer, larynx cancer), lung cancer (e.g., squamous cell carcinoma, small cell (oat cell) carcinoma, large cell carcinoma, and adenocarcinoma), allergic disorders (eosinophilic pneumonia, hypersensitivity pneumonitis (e.g., extrinsic allergic alveolitis, allergic interstitial pneumonitis, organic dust pneumoconiosis, allergic bronchopulmonary aspergillosis, asthma, Wegener's granulomatosis (granulomatous vasculitis), Goodpasture's syndrome)), pneumonia (e.g., bacterial pneumonia (e.g., Streptococcus pneumoniae (pneumoncoccal pneumonia), Staphylococcus aureus (staphylococcal pneumonia), Gram-negative bacterial pneumonia (caused by, e.g., Klebsiella and Pseudomas spp.), Mycoplasma pneumoniae pneumonia, Hemophilus influenzae pneumonia, Legionella pneumophila (Legionnaires' disease), and Chlamydia psittaci (Psittacosis)), and viral pneumonia (e.g., influenza, chickenpox (varicella).

Additional diseases and disorders of the respiratory system include, but are not limited to bronchiolitis, polio (poliomyelitis), croup, respiratory syncytial viral infection, mumps, erythema infectiosum (fifth disease), roseola infantum, progressive rubella panencephalitis, german measles, and subacute sclerosing panencephalitis), fungal pneumonia (e.g., Histoplasmosis, Coccidioidomycosis, Blastomycosis, fungal infections in people with severely suppressed immune systems (e.g., cryptococcosis, caused by *Cryptococcus neoformans*; aspergillosis, caused by *Aspergillus spp.*; candidiasis, caused by *Candida*; and mucormycosis)), *Pneumocystis carinii* (pneumocystis pneumonia), atypical pneumonias (e.g., *Mycoplasma* and *Chlamydia* spp.), opportunistic infection pneumonia, nosocomial pneumonia, chemical

pneumonitis, and aspiration pneumonia, pleural disorders (e.g., pleurisy, pleural effusion, and pneumothorax (e.g., simple spontaneous pneumothorax, complicated spontaneous pneumothorax, tension pneumothorax)), obstructive airway diseases (e.g., asthma, chronic obstructive pulmonary disease (COPD), emphysema, chronic or acute bronchitis), occupational lung diseases (e.g., silicosis, black lung (coal workers' pneumoconiosis), asbestosis, berylliosis, occupational asthsma, byssinosis, and benign pneumoconioses), Infiltrative Lung Disease (e.g., pulmonary fibrosis (e.g., fibrosing alveolitis, usual interstitial pneumonia), idiopathic pulmonary fibrosis, desquamative interstitial pneumonia, lymphoid interstitial pneumonia, histiocytosis X (e.g., Letterer-Siwe disease, Hand-Schüller-Christian disease, eosinophilic granuloma), idiopathic pulmonary hemosiderosis, sarcoidosis and pulmonary alveolar proteinosis), Acute respiratory distress syndrome (also called, e.g., adult respiratory distress syndrome), edema, pulmonary embolism, bronchitis (e.g., viral, bacterial), bronchiectasis, atelectasis, lung abscess (caused by, e.g., *Staphylococcus aureus* or *Legionella pneumophila*), and cystic fibrosis.

Anti-Angiogenesis Activity

The naturally occurring balance between endogenous stimulators and [0665] inhibitors of angiogenesis is one in which inhibitory influences predominate. Rastinejad et al., Cell 56:345-355 (1989). In those rare instances in which neovascularization occurs under normal physiological conditions, such as wound healing, organ regeneration, embryonic development, and female reproductive processes, angiogenesis is stringently regulated and spatially and temporally delimited. Under conditions of pathological angiogenesis such as that characterizing solid tumor growth, these regulatory controls fail. Unregulated angiogenesis becomes pathologic and sustains progression of many neoplastic and non-neoplastic diseases. A number of serious diseases are dominated by abnormal neovascularization including solid tumor growth and metastases, arthritis, some types of eye disorders, and psoriasis. See, e.g., reviews by Moses et al., Biotech. 9:630-634 (1991); Folkman et al., N. Engl. J. Med., 333:1757-1763 (1995); Auerbach et al., J. Microvasc. Res. 29:401-411 (1985); Folkman, Advances in Cancer Research, eds. Klein and Weinhouse, Academic Press, New York, pp. 175-203 (1985); Patz, Am. J. Opthalmol. 94:715-743 (1982); and

Folkman et al., Science 221:719-725 (1983). In a number of pathological conditions, the process of angiogenesis contributes to the disease state. For example, significant data have accumulated which suggest that the growth of solid tumors is dependent on angiogenesis. Folkman and Klagsbrun, *Science* 235:442-447 (1987).

The present invention provides for treatment of diseases or disorders [0666] associated with neovascularization by administration of the polynucleotides and/or polypeptides of the invention, as well as agonists or antagonists of the present invention. Malignant and metastatic conditions which can be treated with the polynucleotides and polypeptides, or agonists or antagonists of the invention include, but are not limited to, malignancies, solid tumors, and cancers described herein and otherwise known in the art (for a review of such disorders, see Fishman et al., Medicine, 2d Ed., J. B. Lippincott Co., Philadelphia (1985)). Thus, the present invention provides a method of treating an angiogenesis-related disease and/or disorder, comprising administration to an individual in need thereof a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist of the invention. For example, polynucleotides, polypeptides, antagonists and/or agonists may be utilized in a variety of additional methods in order to therapeutically treat a cancer or tumor. Cancers which may be treated with polynucleotides, polypeptides, antagonists and/or agonists include, but are not limited to solid tumors, including prostate, lung, breast, ovarian, stomach, pancreas, larynx, esophagus, testes, liver, parotid, biliary tract, colon, rectum, cervix, uterus, endometrium, kidney, bladder, thyroid cancer; primary tumors and metastases; melanomas; glioblastoma; Kaposi's sarcoma; leiomyosarcoma; non- small cell lung cancer; colorectal cancer; advanced malignancies; and blood born tumors such as leukemias. For example, polynucleotides, polypeptides, antagonists and/or agonists may be delivered topically, in order to treat cancers such as skin cancer, head and neck tumors, breast tumors, and Kaposi's sarcoma.

[0667] Within yet other aspects, polynucleotides, polypeptides, antagonists and/or agonists may be utilized to treat superficial forms of bladder cancer by, for example, intravesical administration. Polynucleotides, polypeptides, antagonists and/or agonists may be delivered directly into the tumor, or near the tumor site, via injection or a catheter. Of course, as the artisan of ordinary skill will appreciate, the appropriate

mode of administration will vary according to the cancer to be treated. Other modes of delivery are discussed herein.

Polynucleotides, polypeptides, antagonists and/or agonists may be useful in treating other disorders, besides cancers, which involve angiogenesis. These disorders include, but are not limited to: benign tumors, for example hemangiomas, acoustic neuromas, neurofibromas, trachomas, and pyogenic granulomas; artheroscleric plaques; ocular angiogenic diseases, for example, diabetic retinopathy, retinopathy of prematurity, macular degeneration, corneal graft rejection, neovascular glaucoma, retrolental fibroplasia, rubeosis, retinoblastoma, uvietis and Pterygia (abnormal blood vessel growth) of the eye; rheumatoid arthritis; psoriasis; delayed wound healing; endometriosis; vasculogenesis; granulations; hypertrophic scars (keloids); nonunion fractures; scleroderma; trachoma; vascular adhesions; myocardial angiogenesis; coronary collaterals; cerebral collaterals; arteriovenous malformations; ischemic limb angiogenesis; Osler-Webber Syndrome; plaque neovascularization; telangiectasia; hemophiliac joints; angiofibroma; fibromuscular dysplasia; wound granulation; Crohn's disease; and atherosclerosis.

[0669] For example, within one aspect of the present invention methods are provided for treating hypertrophic scars and keloids, comprising the step of administering a polynucleotide, polypeptide, antagonist and/or agonist of the invention to a hypertrophic scar or keloid.

[0670] Within one embodiment of the present invention polynucleotides, polypeptides, antagonists and/or agonists of the invention are directly injected into a hypertrophic scar or keloid, in order to prevent the progression of these lesions. This therapy is of particular value in the prophylactic treatment of conditions which are known to result in the development of hypertrophic scars and keloids (e.g., burns), and is preferably initiated after the proliferative phase has had time to progress (approximately 14 days after the initial injury), but before hypertrophic scar or keloid development. As noted above, the present invention also provides methods for treating neovascular diseases of the eye, including for example, corneal neovascularization, neovascular glaucoma, proliferative diabetic retinopathy, retrolental fibroplasia and macular degeneration.

[0671] Moreover, ocular disorders associated with neovascularization which can

be treated with the polynucleotides and polypeptides of the present invention (including agonists and/or antagonists) include, but are not limited to: neovascular glaucoma, diabetic retinopathy, retinoblastoma, retrolental fibroplasia, uveitis, retinopathy of prematurity macular degeneration, corneal graft neovascularization, as well as other eye inflammatory diseases, ocular tumors and diseases associated with choroidal or iris neovascularization. See, e.g., reviews by Waltman et al., Am. J. Ophthal. 85:704-710 (1978) and Gartner et al., Surv. Ophthal. 22:291-312 (1978).

Thus, within one aspect of the present invention methods are provided for [0672] treating neovascular diseases of the eye such as comeal neovascularization (including corneal graft neovascularization), comprising the step of administering to a patient a therapeutically effective amount of a compound (as described above) to the comea, such that the formation of blood vessels is inhibited. Briefly, the cornea is a tissue, which normally lacks blood vessels. In certain pathological conditions however, capillaries may extend into the cornea from the pericorneal vascular plexus of the limbus. When the cornea becomes vascularized, it also becomes clouded, resulting in a decline in the patient's visual acuity. Visual loss may become complete if the cornea completely opacitates. A wide variety of disorders can result in corneal neovascularization, including for example, corneal infections (e.g., trachoma, herpes simplex keratitis, leishmaniasis and onchocerciasis), immunological processes (e.g., graft rejection and Stevens-Johnson's syndrome), alkali burns, trauma, inflammation (of any cause), toxic and nutritional deficiency states, and as a complication of wearing contact lenses.

[0673] Within particularly preferred embodiments of the invention, may be prepared for topical administration in saline (combined with any of the preservatives and antimicrobial agents commonly used in ocular preparations), and administered in eyedrop form. The solution or suspension may be prepared in its pure form and administered several times daily. Alternatively, anti-angiogenic compositions; prepared as described above, may also be administered directly to the cornea. Within preferred embodiments, the anti-angiogenic composition is prepared with a muco-adhesive polymer, which binds to cornea. Within further embodiments, the anti-angiogenic factors or anti-angiogenic compositions may be utilized as an adjunct to conventional steroid therapy. Topical therapy may also be useful prophylactically in

comeal lesions which are known to have a high probability of inducing an angiogenic response (such as chemical burns). In these instances the treatment, likely in combination with steroids, may be instituted immediately to help prevent subsequent complications.

injected directly into the corneal stroma by an ophthalmologist under microscopic guidance. The preferred site of injection may vary with the morphology of the individual lesion, but the goal of the administration would be to place the composition at the advancing front of the vasculature (i.e., interspersed between the blood vessels and the normal cornea). In most cases this would involve perilimbic corneal injection to "protect" the cornea from the advancing blood vessels. This method may also be utilized shortly after a corneal insult in order to prophylactically prevent corneal neovascularization. In this situation, the material could be injected in the perilimbic cornea interspersed between the corneal lesion and its undesired potential limbic blood supply. Such methods may also be utilized in a similar fashion to prevent capillary invasion of transplanted corneas. In a sustained-release form, injections might only be required 2-3 times per year. A steroid could also be added to the injection solution to reduce inflammation resulting from the injection itself.

treating neovascular glaucoma, comprising the step of administering to a patient a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist to the eye, such that the formation of blood vessels is inhibited. In one embodiment, the compound may be administered topically to the eye in order to treat early forms of neovascular glaucoma. Within other embodiments, the compound may be implanted by injection into the region of the anterior chamber angle. Within other embodiments, the compound may also be placed in any location such that the compound is continuously released into the aqueous humor. Within another aspect of the present invention, methods are provided for treating proliferative diabetic retinopathy, comprising the step of administering to a patient a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist to the eyes, such that the formation of blood vessels is inhibited.

[0676] Within particularly preferred embodiments of the invention, proliferative

diabetic retinopathy may be treated by injection into the aqueous humor or the vitreous, in order to increase the local concentration of the polynucleotide, polypeptide, antagonist and/or agonist in the retina. Preferably, this treatment should be initiated prior to the acquisition of severe disease requiring photocoagulation.

[0677] Within another aspect of the present invention, methods are provided for treating retrolental fibroplasia, comprising the step of administering to a patient a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist to the eye, such that the formation of blood vessels is inhibited. The compound may be administered topically, via intravitreous injection and/or via intraocular implants.

[0678] Additionally, disorders which can be treated with the polynucleotides, polypeptides, agonists and/or agonists include, but are not limited to, hemangioma, arthritis, psoriasis, angiofibroma, atherosclerotic plaques, delayed wound healing, granulations, hemophilic joints, hypertrophic scars, nonunion fractures, Osler-Weber syndrome, pyogenic granuloma, scleroderma, trachoma, and vascular adhesions.

Moreover, disorders and/or states, which can be treated, prevented, [0679] diagnosed and/or prognosed with the polynucleotides, polypeptides, agonists and/or agonists of the invention include, but are not limited to, solid tumors, blood born tumors such as leukemias, tumor metastasis, Kaposi's sarcoma, benign tumors, for example hemangiomas, acoustic neuromas, neurofibromas, trachomas, and pyogenic granulomas, rheumatoid arthritis, psoriasis, ocular angiogenic diseases, for example, diabetic retinopathy, retinopathy of prematurity, macular degeneration, comeal graft rejection, neovascular glaucoma, retrolental fibroplasia, rubeosis, retinoblastoma, and uvietis, delayed wound healing, endometriosis, vascluogenesis, granulations, hypertrophic scars (keloids), nonunion fractures, scleroderma, trachoma, vascular adhesions, myocardial angiogenesis, coronary collaterals, cerebral collaterals, arteriovenous malformations, ischemic limb angiogenesis, Osler-Webber Syndrome, neovascularization, telangiectasia, hemophiliac joints, fibromuscular dysplasia, wound granulation, Crohn's disease, atherosclerosis, birth control agent by preventing vascularization required for embryo implantation controlling menstruation, diseases that have angiogenesis as a pathologic consequence

such as cat scratch disease (Rochele minalia quintosa), ulcers (Helicobacter pylon), Bartonellosis and bacillary angiomatosis.

ln one aspect of the birth control method, an amount of the compound sufficient to block embryo implantation is administered before or after intercourse and fertilization have occurred, thus providing an effective method of birth control, possibly a "morning after" method. Polynucleotides, polypeptides, agonists and/or agonists may also be used in controlling menstruation or administered as either a peritoneal lavage fluid or for peritoneal implantation in the treatment of endometriosis.

[0681] Polynucleotides, polypeptides, agonists and/or agonists of the present invention may be incorporated into surgical sutures in order to prevent stitch granulomas.

Polynucleotides, polypeptides, agonists and/or agonists may be utilized in a [0682] wide variety of surgical procedures. For example, within one aspect of the present invention a compositions (in the form of, for example, a spray or film) may be utilized to coat or spray an area prior to removal of a tumor, in order to isolate normal surrounding tissues from malignant tissue, and/or to prevent the spread of disease to surrounding tissues. Within other aspects of the present invention, compositions (e.g., in the form of a spray) may be delivered via endoscopic procedures in order to coat tumors, or inhibit angiogenesis in a desired locale. Within yet other aspects of the present invention, surgical meshes, which have been coated with anti- angiogenic compositions of the present invention may be utilized in any procedure wherein a surgical mesh might be utilized. For example, within one embodiment of the invention a surgical mesh laden with an anti-angiogenic composition may be utilized during abdominal cancer resection surgery (e.g., subsequent to colon resection) in order to provide support to the structure, and to release an amount of the antiangiogenic factor.

[0683] Within further aspects of the present invention, methods are provided for treating tumor excision sites, comprising administering a polynucleotide, polypeptide, agonist and/or agonist to the resection margins of a tumor subsequent to excision, such that the local recurrence of cancer and the formation of new blood vessels at the site is inhibited. Within one embodiment of the invention, the anti-angiogenic compound is administered directly to the tumor excision site (e.g., applied by swabbing, brushing or

otherwise coating the resection margins of the tumor with the anti-angiogenic compound). Alternatively, the anti-angiogenic compounds may be incorporated into known surgical pastes prior to administration. Within particularly preferred embodiments of the invention, the anti-angiogenic compounds are applied after hepatic resections for malignancy, and after neurosurgical operations.

[0684] Within one aspect of the present invention, polynucleotides, polypeptides, agonists and/or agonists may be administered to the resection margin of a wide variety of tumors, including for example, breast, colon, brain and hepatic tumors. For example, within one embodiment of the invention, anti-angiogenic compounds may be administered to the site of a neurological tumor subsequent to excision, such that the formation of new blood vessels at the site are inhibited.

[0685] The polynucleotides, polypeptides, agonists and/or agonists of the present invention may also be administered along with other anti-angiogenic factors. Representative examples of other anti-angiogenic factors include: Anti-Invasive Factor, retinoic acid and derivatives thereof, paclitaxel, Suramin, Tissue Inhibitor of Metalloproteinase-1, Tissue Inhibitor of Metalloproteinase-2, Plasminogen Activator Inhibitor-1, Plasminogen Activator Inhibitor-2, and various forms of the lighter "d group" transition metals.

[0686] Lighter "d group" transition metals include, for example, vanadium, molybdenum, tungsten, titanium, niobium, and tantalum species. Such transition metal species may form transition metal complexes. Suitable complexes of the above-mentioned transition metal species include oxo transition metal complexes.

[0687] Representative examples of vanadium complexes include oxo vanadium complexes such as vanadate and vanadyl complexes. Suitable vanadate complexes include metavanadate and orthovanadate complexes such as, for example, ammonium metavanadate, sodium metavanadate, and sodium orthovanadate. Suitable vanadyl complexes include, for example, vanadyl acetylacetonate and vanadyl sulfate including vanadyl sulfate hydrates such as vanadyl sulfate mono- and trihydrates.

[0688] Representative examples of tungsten and molybdenum complexes also include oxo complexes. Suitable oxo tungsten complexes include tungstate and tungsten oxide complexes. Suitable tungstate complexes include ammonium tungstate, calcium tungstate, sodium tungstate dihydrate, and tungstic acid. Suitable

tungsten oxides include tungsten (IV) oxide and tungsten (VI) oxide. Suitable oxo molybdenum complexes include molybdate, molybdenum oxide, and molybdenyl complexes. Suitable molybdate complexes include ammonium molybdate and its hydrates, sodium molybdate and its hydrates, sodium molybdate and its hydrates. Suitable molybdenum oxides include molybdenum (VI) oxide, molybdenum (VI) oxide, and molybdic acid. Suitable molybdenyl complexes include, for example, molybdenyl acetylacetonate. Other suitable tungsten and molybdenum complexes include hydroxo derivatives derived from, for example, glycerol, tartaric acid, and sugars.

A wide variety of other anti-angiogenic factors may also be utilized within [0689] the context of the present invention. Representative examples include platelet factor 4; protamine sulphate; sulphated chitin derivatives (prepared from queen crab shells), (Murata et al., Cancer Res. 51:22-26 (1991)); Sulphated Polysaccharide Peptidoglycan Complex (SP- PG) (the function of this compound may be enhanced by the presence of steroids such as estrogen, and tamoxifen citrate); Staurosporine; modulators of matrix metabolism, including for example, proline analogs, cishydroxyproline, d,L-3,4-dehydroproline, Thiaproline, alpha,alpha-dipyridyl, aminopropionitrile fumarate; 4-propyl-5-(4-pyridinyl)-2(3H)-oxazolone; Methotrexate; Mitoxantrone; Heparin; Interferons; 2 Macroglobulin-serum; ChIMP-3 (Pavloff et al., J. Bio. Chem. 267:17321-17326 (1992)); Chymostatin (Tomkinson et al., Biochem J. 286:475-480 (1992)); Cyclodextrin Tetradecasulfate; Eponemycin; Camptothecin; Fumagillin (Ingber et al., Nature 348:555-557 (1990)); Gold Sodium Thiomalate ("GST"; Matsubara and Ziff, J. Clin. Invest. 79:1440-1446 (1987)); anticollagenase-serum; alpha2-antiplasmin (Holmes et al., J. Biol. Chem. 262(4):1659-1664 (1987)); Bisantrene (National Cancer Institute); Lobenzarit disodium (N-(2)-carboxyphenyl-4chloroanthronilic acid disodium or "CCA"; Takeuchi et al., Agents Actions 36:312-316, 1992); Thalidomide; Angostatic steroid; AGM-1470; carboxynaminolmidazole; and metalloproteinase inhibitors such as BB94.

Musculoskeletal System Disorders

[0690] Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose

disorders of the musculoskeletal system, including but not limited to, disorders of the bone, joints, ligaments, tendons, bursa, muscle, and/or neoplasms and cancers associated with musculoskeletal tissue.

[0691] Diseases or disorders of the bone include, but are not limited to, Albers-Schönberg disease, bowlegs, heel spurs, Köhler's bone disease, knock-knees, Legg-Calvé-Perthes disease, Marfan's syndrome, mucopolysaccharidoses, Osgood-Schlatter disease, osteochondroses, osteochondrodysplasia, osteomyelitis, osteopetroses, osteoporosis (postmenopausal, senile, and juvenile), Paget's disease, Scheuermann's disease, scoliosis, Sever's disease, and patellofemoral stress syndrome.

[0692] Joint diseases or disorders include, but are not limited to, ankylosing spondylitis, Behçet's syndrome, CREST syndrome, Ehlers-Danlos syndrome, infectious arthritis, discoid lupus erythematosus, systemic lupus erythematosus, Lyme disease, osteoarthritis, psoriatic arthritis, relapsing polychondrites, Reiter's syndrome, rheumatoid arthritis (adult and juvenile), scleroderma, and Still's disease.

[0693] Diseases or disorders affecting ligaments, tendons, or bursa include, but are not limited to, ankle sprain, bursitis, posterior Achilles tendon bursitis (Haglund's deformity), anterior Achilles tendon bursitis (Albert's disease), tendinitis, tenosynovitis, poplieus tendinitis, Achilles tendinitis, medial or lateral epicondylitis, rotator cuff tendinitis, spasmodic torticollis, and fibromyalgia syndrome.

[0694] Muscle diseases or disorders include, but are not limited to, Becker's muscular dystrophy, Duchenne's muscular dystrophy, Landouzy-Dejerine muscular dystrophy, Leyden-Möbius muscular dystrophy, Erb's muscular dystrophy, Charcot's joints, dermatomyositis, gout, pseudogout, glycogen storage diseases, Pompe's disease, mitochondrial myopathy, periodic paralysis, polymyalgia rheumatica, polymyositis, Steinert's disease, Thomsen's disease, anterolateral and posteromedial shin splints, posterior femoral muscle strain, and fibromyositis.

[0695] Musculoskeletal tissue may also develop cancers and/or neoplasms that include, but are not limited to, osteochondroma, benign chondroma, chondroblastoma, chondromyxoid fibroma, osteoid osteoma, giant cell tumor, multiple myeloma, osteosarcoma, fibrosarcoma, malignant fibrous histiocytoma, chondrosarcoma, Ewing's tumor, and malignant lymphoma of bone.

Neural Activity and Neurological Diseases

[0696] The polynucleotides, polypeptides and agonists or antagonists of the invention may be used for the diagnosis and/or treatment of diseases, disorders, damage or injury of the brain and/or nervous system. Nervous system disorders that can be treated with the compositions of the invention (e.g., polypeptides, polynucleotides, and/or agonists or antagonists), include, but are not limited to, nervous system injuries, and diseases or disorders which result in either a disconnection of axons, a diminution or degeneration of neurons, or demyelination. Nervous system lesions which may be treated in a patient (including human and nonhuman mammalian patients) according to the methods of the invention, include but are not limited to, the following lesions of either the central (including spinal cord, brain) or peripheral nervous systems: (1) ischemic lesions, in which a lack of oxygen in a portion of the nervous system results in neuronal injury or death, including cerebral infarction or ischemia, or spinal cord infarction or ischemia; (2) traumatic lesions, including lesions caused by physical injury or associated with surgery, for example, lesions which sever a portion of the nervous system, or compression injuries; (3) malignant lesions, in which a portion of the nervous system is destroyed or injured by malignant tissue which is either a nervous system associated malignancy or a malignancy derived from non-nervous system tissue; (4) infectious lesions, in which a portion of the nervous system is destroyed or injured as a result of infection, for example, by an abscess or associated with infection by human immunodeficiency virus, herpes zoster, or herpes simplex virus or with Lyme disease, tuberculosis, or syphilis; (5) degenerative lesions, in which a portion of the nervous system is destroyed or injured as a result of a degenerative process including but not limited to, degeneration associated with Parkinson's disease, Alzheimer's disease, Huntington's chorea, or amyotrophic lateral sclerosis (ALS); (6) lesions associated with nutritional diseases or disorders, in which a portion of the nervous system is destroyed or injured by a nutritional disorder or disorder of metabolism including, but not limited to, vitamin B12 deficiency, folic acid deficiency, Wernicke disease, tobacco-alcohol amblyopia, Marchiafava-Bignami disease (primary degeneration of the corpus callosum), and alcoholic cerebellar degeneration; (7) neurological lesions associated with systemic diseases including, but not limited to, diabetes (diabetic neuropathy,

Bell's palsy), systemic lupus erythematosus, carcinoma, or sarcoidosis; (8) lesions caused by toxic substances including alcohol, lead, or particular neurotoxins; and (9) demyelinated lesions in which a portion of the nervous system is destroyed or injured by a demyelinating disease including, but not limited to, multiple sclerosis, human immunodeficiency virus-associated myelopathy, transverse myelopathy or various etiologies, progressive multifocal leukoencephalopathy, and central pontine myelinolysis.

In one embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to protect neural cells from the damaging effects of hypoxia. In a further preferred embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to protect neural cells from the damaging effects of cerebral hypoxia. According to this embodiment, the compositions of the invention are used to treat or prevent neural cell injury associated with cerebral hypoxia. In one non-exclusive aspect of this embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention, are used to treat or prevent neural cell injury associated with cerebral ischemia. In another non-exclusive aspect of this embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent neural cell injury associated with cerebral infarction.

[0698] In another preferred embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent neural cell injury associated with a stroke. In a specific embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent cerebral neural cell injury associated with a stroke.

[0699] In another preferred embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent neural cell injury associated with a heart attack. In a specific embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent cerebral neural cell injury associated with a heart attack.

[0700] The compositions of the invention which are useful for treating or preventing a nervous system disorder may be selected by testing for biological activity in promoting the survival or differentiation of neurons. For example, and not by way

of limitation, compositions of the invention which elicit any of the following effects may be useful according to the invention: (1) increased survival time of neurons in culture either in the presence or absence of hypoxia or hypoxic conditions; (2) increased sprouting of neurons in culture or in vivo; (3) increased production of a neuron-associated molecule in culture or in vivo, e.g., choline acetyltransferase or acetylcholinesterase with respect to motor neurons; or (4) decreased symptoms of neuron dysfunction in vivo. Such effects may be measured by any method known in the art. In preferred, non-limiting embodiments, increased survival of neurons may routinely be measured using a method set forth herein or otherwise known in the art, such as, for example, in Zhang et al., Proc Natl Acad Sci USA 97:3637-42 (2000) or in Arakawa et al., J. Neurosci., 10:3507-15 (1990); increased sprouting of neurons may be detected by methods known in the art, such as, for example, the methods set forth in Pestronk et al., Exp. Neurol., 70:65-82 (1980), or Brown et al., Ann. Rev. Neurosci., 4:17-42 (1981); increased production of neuron-associated molecules may be measured by bioassay, enzymatic assay, antibody binding, Northern blot assay, etc., using techniques known in the art and depending on the molecule to be measured; and motor neuron dysfunction may be measured by assessing the physical manifestation of motor neuron disorder, e.g., weakness, motor neuron conduction velocity, or functional disability.

[0701] In specific embodiments, motor neuron disorders that may be treated according to the invention include, but are not limited to, disorders such as infarction, infection, exposure to toxin, trauma, surgical damage, degenerative disease or malignancy that may affect motor neurons as well as other components of the nervous system, as well as disorders that selectively affect neurons such as amyotrophic lateral sclerosis, and including, but not limited to, progressive spinal muscular atrophy, progressive bulbar palsy, primary lateral sclerosis, infantile and juvenile muscular atrophy, progressive bulbar paralysis of childhood (Fazio-Londe syndrome), poliomyelitis and the post polio syndrome, and Hereditary Motorsensory Neuropathy (Charcot-Marie-Tooth Disease).

[0702] Further, polypeptides or polynucleotides of the invention may play a role in neuronal survival; synapse formation; conductance; neural differentiation, etc. Thus, compositions of the invention (including polynucleotides, polypeptides, and agonists

or antagonists) may be used to diagnose and/or treat or prevent diseases or disorders associated with these roles, including, but not limited to, learning and/or cognition disorders. The compositions of the invention may also be useful in the treatment or prevention of neurodegenerative disease states and/or behavioural disorders. Such neurodegenerative disease states and/or behavioral disorders include, but are not limited to, Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, Tourette Syndrome, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder, panic disorder, learning disabilities, ALS, psychoses, autism, and altered behaviors, including disorders in feeding, sleep patterns, balance, and perception. In addition, compositions of the invention may also play a role in the treatment, prevention and/or detection of developmental disorders associated with the developing embryo, or sexually-linked disorders.

[0703] Additionally, polypeptides, polynucleotides and/or agonists or antagonists of the invention, may be useful in protecting neural cells from diseases, damage, disorders, or injury, associated with cerebrovascular disorders including, but not limited to, carotid artery diseases (e.g., carotid artery thrombosis, carotid stenosis, or Moyamoya Disease), cerebral amyloid angiopathy, cerebral aneurysm, cerebral anoxia, cerebral arteriosclerosis, cerebral arteriovenous malformations, cerebral artery diseases, cerebral embolism and thrombosis (e.g., carotid artery thrombosis, sinus thrombosis, or Wallenberg's Syndrome), cerebral hemorrhage (e.g., epidural or subdural hematoma, or subarachnoid hemorrhage), cerebral infarction, cerebral ischemia (e.g., transient cerebral ischemia, Subclavian Steal Syndrome, or vertebrobasilar insufficiency), vascular dementia (e.g., multi-infarct), leukomalacia, periventricular, and vascular headache (e.g., cluster headache or migraines).

In accordance with yet a further aspect of the present invention, there is provided a process for utilizing polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, for therapeutic purposes, for example, to stimulate neurological cell proliferation and/or differentiation. Therefore, polynucleotides, polypeptides, agonists and/or antagonists of the invention may be used to treat and/or detect neurologic diseases. Moreover, polynucleotides or polypeptides, or agonists or antagonists of the invention, can be used as a marker or detector of a particular nervous system disease or disorder.

Examples of neurologic diseases which can be treated or detected with [0705] polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include brain diseases, such as metabolic brain diseases which includes phenylketonuria such as maternal phenylketonuria, pyruvate carboxylase deficiency, pyruvate dehydrogenase complex deficiency, Wernicke's Encephalopathy, brain edema, brain neoplasms such as cerebellar neoplasms which include infratentorial neoplasms, cerebral ventricle neoplasms such as choroid plexus neoplasms, hypothalamic neoplasms, supratentorial neoplasms, canavan disease, cerebellar diseases such as cerebellar ataxia which include spinocerebellar degeneration such as ataxia telangiectasia, cerebellar dyssynergia, Friederich's Ataxia, Machado-Joseph Disease, olivopontocerebellar atrophy, cerebellar neoplasms such as infratentorial neoplasms, diffuse cerebral sclerosis such as encephalitis periaxialis, globoid cell leukodystrophy, metachromatic leukodystrophy and subacute panencephalitis.

[0706] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include cerebrovascular disorders (such as carotid artery diseases which include carotid artery thrombosis, carotid stenosis and Moyamoya Disease), cerebral amyloid angiopathy, cerebral aneurysm, cerebral anoxia, cerebral arteriosclerosis, cerebral arteriovenous malformations, cerebral artery diseases, cerebral embolism and thrombosis such as carotid artery thrombosis, sinus thrombosis and Wallenberg's Syndrome, cerebral hemorrhage such as epidural hematoma, subdural hematoma and subarachnoid hemorrhage, cerebral infarction, cerebral ischemia such as transient cerebral ischemia, Subclavian Steal Syndrome and vertebrobasilar insufficiency, vascular dementia such as multi-infarct dementia, periventricular leukomalacia, vascular headache such as cluster headache and migraine.

[0707] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include dementia such as AIDS Dementia Complex, presentile dementia such as Alzheimer's Disease and Creutzfeldt-Jakob Syndrome, senile dementia such as Alzheimer's Disease and progressive supranuclear palsy, vascular dementia such as multi-infarct dementia, encephalitis which include encephalitis periaxialis, viral

encephalitis such as epidemic encephalitis, Japanese Encephalitis, St. Louis Encephalitis, tick-borne encephalitis and West Nile Fever, acute disseminated encephalomyelitis, meningoencephalitis such as uveomeningoencephalitic syndrome, Postencephalitic Parkinson Disease and subacute sclerosing panencephalitis, encephalomalacia such as periventricular leukomalacia, epilepsy such as generalized epilepsy which includes infantile spasms, absence epilepsy, myoclonic epilepsy which includes MERRF Syndrome, tonic-clonic epilepsy, partial epilepsy such as complex partial epilepsy, frontal lobe epilepsy and temporal lobe epilepsy, post-traumatic epilepsy, status epilepticus such as Epilepsia Partialis Continua, and Hallervorden-Spatz Syndrome.

[0708] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include hydrocephalus such as Dandy-Walker Syndrome and normal pressure hydrocephalus, hypothalamic diseases such as hypothalamic neoplasms, cerebral malaria, narcolepsy which includes cataplexy, bulbar poliomyelitis, cerebri pseudotumor, Rett Syndrome, Reye's Syndrome, thalamic diseases, cerebral toxoplasmosis, intracranial tuberculoma and Zellweger Syndrome, central nervous system infections such as AIDS Dementia Complex, Brain Abscess, subdural empyema, encephalomyelitis such as Equine Encephalomyelitis, Venezuelan Equine Encephalomyelitis, Necrotizing Hemorrhagic Encephalomyelitis, Visna, and cerebral malaria.

[0709] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include meningitis such as arachnoiditis, aseptic meningitis such as viral meningitis which includes lymphocytic choriomeningitis, Bacterial meningitis which includes Haemophilus Meningitis, Listeria Meningitis, Meningococcal Meningitis such as Waterhouse-Friderichsen Syndrome, Pneumococcal Meningitis and meningeal tuberculosis, fungal meningitis such as Cryptococcal Meningitis, subdural effusion, meningoencephalitis such as uvemeningoencephalitic syndrome, myelitis such as transverse myelitis, neurosyphilis such as tabes dorsalis, poliomyelitis which includes bulbar poliomyelitis and postpoliomyelitis syndrome, prion diseases (such as

Creutzfeldt-Jakob Syndrome, Bovine Spongiform Encephalopathy, Gerstmann-Straussler Syndrome, Kuru, Scrapie), and cerebral toxoplasmosis.

[0710] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include central nervous system neoplasms such as brain neoplasms that include cerebellar neoplasms such as infratentorial neoplasms, cerebral ventricle neoplasms such as choroid plexus neoplasms, hypothalamic neoplasms and supratentorial neoplasms, meningeal neoplasms, spinal cord neoplasms which include epidural neoplasms, demyelinating diseases such as Canavan Diseases, diffuse cerebral sceloris which includes adrenoleukodystrophy, encephalitis periaxialis, globoid cell leukodystrophy, diffuse cerebral sclerosis such as metachromatic leukodystrophy, allergic encephalomyelitis, necrotizing hemorrhagic encephalomyelitis, progressive multifocal leukoencephalopathy, multiple sclerosis, central pontine myelinolysis, transverse myelitis, neuromyelitis optica, Scrapie, Swayback, Chronic Fatigue Syndrome, Visna, High Pressure Nervous Syndrome, Meningism, spinal cord diseases such as amyotonia congenita, amyotrophic lateral sclerosis, spinal muscular atrophy such as Werdnig-Hoffmann Disease, spinal cord compression, spinal cord neoplasms such as epidural neoplasms, syringomyelia, Tabes Dorsalis, Stiff-Man Syndrome, mental retardation such as Angelman Syndrome, Cri-du-Chat Syndrome, De Lange's Syndrome, Down Syndrome, Gangliosidoses such as gangliosidoses G(M1), Sandhoff Disease, Tay-Sachs Disease, Hartnup Disease, homocystinuria, Laurence-Moon-Biedl Syndrome, Lesch-Nyhan Syndrome, Maple Syrup Urine Disease, mucolipidosis such neuronal ceroid-lipofuscinosis, oculocerebrorenal syndrome, fucosidosis, phenylketonuria such as maternal phenylketonuria, Prader-Willi Syndrome, Rett Syndrome, Rubinstein-Taybi Syndrome, Tuberous Sclerosis, WAGR Syndrome, nervous system abnormalities such as holoprosencephaly, neural tube defects such as anencephaly which includes hydrangencephaly, Arnold-Chairi Deformity. encephalocele, meningocele, meningomyelocele, spinal dysraphism such as spina bifida cystica and spina bifida occulta.

[0711] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include hereditary motor and sensory neuropathies which include Charcot-Marie

Disease, Hereditary optic atrophy, Refsum's Disease, hereditary spastic paraplegia, Werdnig-Hoffmann Disease, Hereditary Sensory and Autonomic Neuropathies such as Congenital Analgesia and Familial Dysautonomia, Neurologic manifestations (such as agnosia that include Gerstmann's Syndrome, Amnesia such as retrograde amnesia, apraxia, neurogenic bladder, cataplexy, communicative disorders such as hearing disorders that includes deafness, partial hearing loss, loudness recruitment and tinnitus, language disorders such as aphasia which include agraphia, anomia, broca aphasia, and Wernicke Aphasia, Dyslexia such as Acquired Dyslexia, language development disorders, speech disorders such as aphasia which includes anomia, broca aphasia and Wernicke Aphasia, articulation disorders, communicative disorders such as speech disorders which include dysarthria, echolalia, mutism and stuttering, voice disorders such as aphonia and hoarseness, decerebrate state, delirium, fasciculation, hallucinations, meningism, movement disorders such as angelman syndrome, ataxia, athetosis, chorea, dystonia, hypokinesia, muscle hypotonia, myoclonus, tic, torticollis and tremor, muscle hypertonia such as muscle rigidity such as stiff-man syndrome, muscle spasticity, paralysis such as facial paralysis which includes Herpes Zoster Oticus, Gastroparesis, Hemiplegia, ophthalmoplegia such as diplopia, Duane's Syndrome, Horner's Syndrome, Chronic progressive external ophthalmoplegia such as Kearns Syndrome, Bulbar Paralysis, Tropical Spastic Paraparesis, Paraplegia such as Brown-Sequard Syndrome, quadriplegia, respiratory paralysis and vocal cord paralysis, paresis, phantom limb, taste disorders such as ageusia and dysgeusia, vision disorders such as amblyopia, blindness, color vision defects, diplopia, hemianopsia, scotoma and subnormal vision, sleep disorders such as hypersomnia which includes Kleine-Levin Syndrome, insomnia, and somnambulism, spasm such as trismus, unconsciousness such as coma, persistent vegetative state and syncope and vertigo, neuromuscular diseases such as amyotonia congenita, amyotrophic lateral sclerosis, Lambert-Eaton Myasthenic Syndrome, motor neuron disease, muscular atrophy such as spinal muscular atrophy, Charcot-Marie Disease and Werdnig-Hoffmann Disease, Postpoliomyelitis Syndrome, Muscular Dystrophy, Myasthenia Gravis, Myotonia Atrophica, Myotonia Confenita, Nemaline Myopathy, Familial Periodic Paralysis, Multiplex Paramyloclonus, Tropical Spastic Paraparesis and Stiff-Man Syndrome, peripheral nervous system diseases such as acrodynia, amyloid neuropathies,

autonomic nervous system diseases such as Adie's Syndrome, Barre-Lieou Syndrome, Familial Dysautonomia, Horner's Syndrome, Reflex Sympathetic Dystrophy and Shy-Drager Syndrome, Cranial Nerve Diseases such as Acoustic Nerve Diseases such as Acoustic Neuroma which includes Neurofibromatosis 2, Facial Nerve Diseases such as Facial Neuralgia, Melkersson-Rosenthal Syndrome, ocular motility disorders which includes amblyopia, nystagmus, oculomotor nerve paralysis, ophthalmoplegia such as Duane's Syndrome, Horner's Syndrome, Chronic Progressive External Ophthalmoplegia which includes Kearns Syndrome, Strabismus such as Esotropia and Exotropia, Oculomotor Nerve Paralysis, Optic Nerve Diseases such as Optic Atrophy which includes Hereditary Optic Atrophy, Optic Disk Drusen, Optic Neuritis such as Neuromyelitis Optica, Papilledema, Trigeminal Neuralgia, Vocal Cord Paralysis, Demyelinating Diseases such as Neuromyelitis Optica and Swayback, and Diabetic neuropathies such as diabetic foot.

[0712] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include nerve compression syndromes such as carpal tunnel syndrome, tarsal tunnel syndrome, thoracic outlet syndrome such as cervical rib syndrome, ulnar nerve compression syndrome, neuralgia such as causalgia, cervico-brachial neuralgia, facial neuralgia and trigeminal neuralgia, neuritis such as experimental allergic neuritis, optic neuritis, polyneuritis, polyradiculoneuritis and radiculities such as polyradiculitis, hereditary motor and sensory neuropathies such as Charcot-Marie Disease, Hereditary Optic Atrophy, Refsum's Disease, Hereditary Spastic Paraplegia and Werdnig-Hoffmann Disease, Hereditary Sensory and Autonomic Neuropathies which include Congenital Analgesia and Familial Dysautonomia, POEMS Syndrome, Sciatica, Gustatory Sweating and Tetany).

Endocrine Disorders

[0713] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose disorders and/or diseases related to hormone imbalance, and/or disorders or diseases of the endocrine system.

growth, sexual function, metabolism, and other functions. Disorders may be classified in two ways: disturbances in the production of hormones, and the inability of tissues to respond to hormones. The etiology of these hormone imbalance or endocrine system diseases, disorders or conditions may be genetic, somatic, such as cancer and some autoimmune diseases, acquired (e.g., by chemotherapy, injury or toxins), or infectious. Moreover, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention can be used as a marker or detector of a particular disease or disorder related to the endocrine system and/or hormone imbalance.

[0715] Endocrine system and/or hormone imbalance and/or diseases encompass disorders of uterine motility including, but not limited to: complications with pregnancy and labor (e.g., pre-term labor, post-term pregnancy, spontaneous abortion, and slow or stopped labor); and disorders and/or diseases of the menstrual cycle (e.g., dysmenorrhea and endometriosis).

[0716] Endocrine system and/or hormone imbalance disorders and/or diseases include disorders and/or diseases of the pancreas, such as, for example, diabetes mellitus, diabetes insipidus, congenital pancreatic agenesis, pheochromocytoma--islet cell tumor syndrome; disorders and/or diseases of the adrenal glands such as, for example, Addison's Disease, corticosteroid deficiency, virilizing disease, hirsutism, Cushing's Syndrome, hyperaldosteronism, pheochromocytoma; disorders and/or diseases of the pituitary gland, such as, for example, hyperpituitarism, hypopituitarism, pituitary dwarfism, pituitary adenoma, panhypopituitarism, acromegaly, gigantism; disorders and/or diseases of the thyroid, including but not limited to, hyperthyroidism, hypothyroidism, Plummer's disease, Graves' disease (toxic diffuse goiter), toxic nodular goiter, thyroiditis (Hashimoto's thyroiditis, subacute granulomatous thyroiditis, and silent lymphocytic thyroiditis), Pendred's syndrome, myxedema, cretinism, thyrotoxicosis, thyroid hormone coupling defect, thymic aplasia, Hurthle cell tumours of the thyroid, thyroid cancer, thyroid carcinoma, Medullary thyroid carcinoma; disorders and/or diseases of the parathyroid, such as, for example, hyperparathyroidism, hypoparathyroidism; disorders and/or diseases hypothalamus.

[0717] In addition, endocrine system and/or hormone imbalance disorders and/or diseases may also include disorders and/or diseases of the testes or ovaries, including cancer. Other disorders and/or diseases of the testes or ovaries further include, for example, ovarian cancer, polycystic ovary syndrome, Klinefelter's syndrome, vanishing testes syndrome (bilateral anorchia), congenital absence of Leydig's cells, cryptorchidism, Noonan's syndrome, myotonic dystrophy, capillary haemangioma of the testis (benign), neoplasias of the testis and neo-testis.

[0718] Moreover, endocrine system and/or hormone imbalance disorders and/or diseases may also include disorders and/or diseases such as, for example, polyglandular deficiency syndromes, pheochromocytoma, neuroblastoma, multiple Endocrine neoplasia, and disorders and/or cancers of endocrine tissues.

[0719] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to diagnose, prognose, prevent, and/or treat endocrine diseases and/or disorders associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1A, column 7 (Tissue Distribution Library Code).

Gastrointestinal Disorders

[0720] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose gastrointestinal disorders, including inflammatory diseases and/or conditions, infections, cancers (e.g., intestinal neoplasms (carcinoid tumor of the small intestine, non-Hodgkin's lymphoma of the small intestine, small bowl lymphoma), and ulcers, such as peptic ulcers.

[0721] Gastrointestinal disorders include dysphagia, odynophagia, inflammation of the esophagus, peptic esophagitis, gastric reflux, submucosal fibrosis and stricturing, Mallory-Weiss lesions, leiomyomas, lipomas, epidermal cancers, adeoncarcinomas, gastric retention disorders, gastroenteritis, gastric atrophy, gastric/stomach cancers, polyps of the stomach, autoimmune disorders such as pernicious anemia, pyloric stenosis, gastritis (bacterial, viral, eosinophilic, stressinduced, chronic erosive, atrophic, plasma cell, and Ménétrier's), and peritoneal diseases (e.g., chyloperioneum, hemoperitoneum, mesenteric cyst, mesenteric

lymphadenitis, mesenteric vascular occlusion, panniculitis, neoplasms, peritonitis, pneumoperitoneum, bubphrenic abscess).

intestine, such as malabsorption syndromes, distension, irritable bowel syndrome, sugar intolerance, celiac disease, duodenal ulcers, duodenitis, tropical sprue. Whipple's disease, intestinal lymphangiectasia, Crohn's disease, appendicitis, obstructions of the ileum, Meckel's diverticulum, multiple diverticula, failure of complete rotation of the small and large intestine, lymphoma, and bacterial and parasitic diseases (such as Traveler's diarrhea, typhoid and paratyphoid, cholera, infection by Roundworms (Ascariasis lumbricoides), Hookworms (Ancylostoma duodenale), Threadworms (Enterobius vermicularis), Tapeworms (Taenia saginata, Echinococcus granulosus, Diphyllobothrium spp., and T. solium).

[0723] Liver diseases and/or disorders include intrahepatic cholestasis (alagille syndrome, biliary liver cirrhosis), fatty liver (alcoholic fatty liver, reye syndrome), hepatic vein thrombosis, hepatolentricular degeneration. hepatomegaly, hepatopulmonary syndrome, hepatorenal syndrome, portal hypertension (esophageal and gastric varices), liver abscess (amebic liver abscess), liver cirrhosis (alcoholic, biliary and experimental), alcoholic liver diseases (fatty liver, hepatitis, cirrhosis), parasitic (hepatic echinococcosis, fascioliasis, amebic liver abscess), jaundice (hemolytic, hepatocellular, and cholestatic), cholestasis, portal hypertension, liver enlargement, ascites, hepatitis (alcoholic hepatitis, animal hepatitis, chronic hepatitis (autoimmune, hepatitis B, hepatitis C, hepatitis D, drug induced), toxic hepatitis, viral human hepatitis (hepatitis A, hepatitis B, hepatitis C, hepatitis D, hepatitis E), Wilson's disease, granulomatous hepatitis, secondary biliary cirrhosis, hepatic encephalopathy, portal hypertension, varices, hepatic encephalopathy, primary biliary cirrhosis, primary sclerosing cholangitis, hepatocellular adenoma, hemangiomas, bile stones, liver failure (hepatic encephalopathy, acute liver failure), and liver neoplasms (angiomyolipoma, calcified liver metastases, cystic liver metastases, epithelial tumors, fibrolamellar hepatocarcinoma, focal nodular hyperplasia, hepatic adenoma, hepatobiliary cystadenoma, hepatoblastoma, hepatocellular carcinoma, hepatoma, liver cancer, liver hemangioendothelioma, mesenchymal hamartoma, mesenchymal tumors of liver, nodular regenerative hyperplasia, benign liver tumors (Hepatic cysts [Simple

cysts, Polycystic liver disease, Hepatobiliary cystadenoma, Choledochal cyst], Mesenchymal tumors [Mesenchymal hamartoma, Infantile hemangioendothelioma, Hemangioma, Peliosis hepatis, Lipomas, Inflammatory pseudotumor, Miscellaneous], Epithelial tumors [Bile duct epithelium (Bile duct hamartoma, Bile duct adenoma), Hepatocyte (Adenoma, Focal nodular hyperplasia, Nodular regenerative hyperplasia)], malignant liver tumors [hepatocellular, hepatoblastoma, hepatocellular carcinoma, cholangiocellular, cholangiocarcinoma, cystadenocarcinoma, tumors of blood vessels, angiosarcoma, Karposi's sarcoma, hemangioendothelioma, other tumors, embryonal sarcoma, fibrosarcoma, leiomyosarcoma, rhabdomyosarcoma, carcinosarcoma, teratoma, carcinoid, squamous carcinoma, primary lymphoma]), peliosis hepatis, erythrohepatic porphyria, hepatic porphyria (acute intermittent porphyria, porphyria cutanea tarda), Zellweger syndrome).

[0724] Pancreatic diseases and/or disorders include acute pancreatitis, chronic pancreatitis (acute necrotizing pancreatitis, alcoholic pancreatitis), neoplasms (adenocarcinoma of the pancreas, cystadenocarcinoma, insulinoma, gastrinoma, and glucagonoma, cystic neoplasms, islet-cell tumors, pancreoblastoma), and other pancreatic diseases (e.g., cystic fibrosis, cyst (pancreatic pseudocyst, pancreatic fistula, insufficiency)).

[0725] Gallbladder diseases include gallstones (cholelithiasis and choledocholithiasis), postcholecystectomy syndrome, diverticulosis of the gallbladder, acute cholecystitis, chronic cholecystitis, bile duct tumors, and mucocele.

Diseases and/or disorders of the large intestine include antibiotic-associated colitis, diverticulitis, ulcerative colitis, acquired megacolon, abscesses, fungal and bacterial infections, anorectal disorders (e.g., fissures, hemorrhoids), colonic diseases (colitis, colonic neoplasms [colon cancer, adenomatous colon polyps (e.g., villous adenoma), colon carcinoma, colorectal cancer], colonic diverticulitis, colonic diverticulosis, megacolon [Hirschsprung disease, toxic megacolon]; sigmoid diseases [proctocolitis, sigmoin neoplasms]), constipation, Crohn's disease, diarrhea (infantile diarrhea, dysentery), duodenal diseases (duodenal neoplasms, duodenal obstruction, duodenal ulcer, duodenitis), enteritis (enterocolitis), HIV enteropathy, ileal diseases (ileal neoplasms, ileitis), immunoproliferative small intestinal disease, inflammatory bowel disease (ulcerative colitis, Crohn's disease), intestinal atresia, parasitic diseases

(anisakiasis, balantidiasis, blastocystis infections, cryptosporidiosis, dientamoebiasis, amebic dysentery, giardiasis), intestinal fistula (rectal fistula), intestinal neoplasms (cecal neoplasms, colonic neoplasms, duodenal neoplasms, ileal neoplasms, intestinal polyps, jejunal neoplasms, rectal neoplasms), intestinal obstruction (afferent loop syndrome, duodenal obstruction, impacted feces, intestinal pseudo-obstruction [cecal volvulus], intussusception), intestinal perforation, intestinal polyps (colonic polyps, gardner syndrome, peutz-jeghers syndrome), jejunal diseases (jejunal neoplasms), malabsorption syndromes (blind loop syndrome, celiac disease, lactose intolerance, short bowl syndrome, tropical sprue, whipple's disease), mesenteric vascular occlusion, pneumatosis cystoides intestinalis, protein-losing enteropathies (intestinal lymphagiectasis), rectal diseases (anus diseases, fecal incontinence, hemorrhoids, proctitis, rectal fistula, rectal prolapse, rectocele), peptic ulcer (duodenal ulcer, peptic esophagitis, hemorrhage, perforation, stomach ulcer, Zollinger-Ellison syndrome), postgastrectomy syndromes (dumping syndrome), stomach diseases (e.g., achlorhydria, duodenogastric reflux (bile reflux), gastric antral vascular ectasia, gastric fistula, gastric outlet obstruction, gastritis (atrophic or hypertrophic), gastroparesis, stomach dilatation, stomach diverticulum, stomach neoplasms (gastric cancer, gastric polyps, gastric adenocarcinoma, hyperplastic gastric polyp), stomach rupture, stomach ulcer, stomach volvulus), tuberculosis, visceroptosis, vomiting (e.g., hematemesis, hyperemesis gravidarum, postoperative nausea and vomiting) and hemorrhagic colitis.

Further diseases and/or disorders of the gastrointestinal system include biliary tract diseases, such as, gastroschisis, fistula (e.g., biliary fistula, esophageal fistula, gastric fistula, intestinal fistula, pancreatic fistula), neoplasms (e.g., biliary tract neoplasms, esophageal neoplasms, such as adenocarcinoma of the esophagus, esophageal squamous cell carcinoma, gastrointestinal neoplasms, pancreatic neoplasms, such as adenocarcinoma of the pancreas, mucinous cystic neoplasm of the pancreas, pancreatic cystic neoplasms, pancreatoblastoma, and peritoneal neoplasms), esophageal disease (e.g., bullous diseases, candidiasis, glycogenic acanthosis, ulceration, barrett esophagus varices, atresia, cyst, diverticulum (e.g., Zenker's diverticulum), fistula (e.g., tracheoesophageal fistula), motility disorders (e.g., CREST syndrome, deglutition disorders, achalasia, spasm, gastroesophageal reflux), neoplasms, perforation (e.g., Boerhaave syndrome, Mallory-Weiss syndrome),

stenosis, esophagitis, diaphragmatic hernia (e.g., hiatal hernia); gastrointestinal diseases, such as, gastroenteritis (e.g., cholera morbus, norwalk virus infection), hemorrhage (e.g., hematemesis, melena, peptic ulcer hemorrhage), stomach neoplasms (gastric cancer, gastric polyps, gastric adenocarcinoma, stomach cancer)), hernia (e.g., congenital diaphragmatic hernia, femoral hernia, inguinal hernia, obturator hernia, umbilical hernia, ventral hernia), and intestinal diseases (e.g., cecal diseases (appendicitis, cecal neoplasms)).

Developmental and Inherited Disorders

[0728] Polynuceotides or polypeptides, or agonists or antagonists of the present invention may be used to treat, prevent, diagnose, and/or prognose diseases associated with mixed fetal tissues, including, but not limited to, developmental and inherited disorders or defects of the nervous system, musculoskelelal system, execretory system, cardiovascular system, hematopoietic system, gastrointestinal system, reproductive system, and respiratory system. Compositions of the present invention may also be used to treat, prevent, diagnose, and/or prognose developmental and inherited disorders or defects associated with, but not limited to, skin, hair, visual, and auditory tissues, metabolism. Additionally, the compositions of the invention may be useful in the diagnosis, treatment, and/or prevention of disorders or diseases associated with, but not limited to, chromosomal or genetic abnormalities and hyperproliferation or neoplasia.

Disorders or defects of the nervous system associated with developmental or inherited abnormalities that may be diagnosed, treated, and/or prevented with the compostions of the invention include, but are not limited to, adrenoleukodystrophy, agenesis of corpus callosum, Alexander disease, anencephaly, Angelman syndrome, Amold-Chiari deformity, Batten disease, Canavan disease, cephalic disorders, Charcot-Marie-Tooth disease, encephalocele, Friedreich's ataxia, Gaucher's disease, Gorlin syndrome, Hallervorden-Spatz disease, hereditary spastic paraplegia, Huntington disease, hydranencephaly, hydrocephalus, Joubert syndrome, Lesch-Nyhan syndrome, leukodystrophy, Menkes disease, microcephaly, Niemann-Pick Type C1, neurofibromatosis, porencephaly, progeria, proteus syndrome, Refsum

disease, spina bifida, Sturge-Weber syndrome, Tay-Sachs disease, tuberous sclerosis, and von Hippel-Lindau disease.

the musculoskeletal system that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, achondroplasia, atlanto-occipital fusion, arthrogryposis mulitplex congenita, autosomal recessive muscular dystrophy, Becker's muscular dystrophy, cerebral palsy, choanal atresia, cleft lip, cleft palate, clubfoot, congenital amputation, congenital dislocation of the hip, congenital torticollis, congenital scoliosis, dopa-repsonsive dystonia, Duchenne muscular dystrophy, early-onset generalized dystonia, femoral torsion, Gorlin syndrome, hypophosphatasia, Klippel-Feil syndrome, knee dislocation, myoclonic dystonia, myotonic dystrophy, nail-patella syndrome, osteogenesis imperfecta, paroxysmal dystonia, progeria, prune-belly syndrome, rapid-onset dystonia parkinsonism, scolosis, syndactyly, Treacher Collins' syndrome, velocardiofacial syndrome, and X-linked dystonia-parkinsonism.

that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, Alport's syndrome, Bartter's syndrome, bladder diverticula, bladder exstrophy, cystinuria, epispadias, Fanconi's syndrome, Hartnup disease, horseshoe kidney, hypospadias, kidney agenesis, kidney ectopia, kidney malrotation, Liddle's syndrome, medullary cystic disease, medullary sponge, multicystic kidney, kidney polycystic kidney disease, nail-patella syndrome, Potter's syndrome, urinary tract flow obstruction, vitamin D-resistant rickets, and Wilm's tumor.

[0732] Cardiovascular disorders or defects of developmental or hereditary origin that may be diagnosed, treated, and/or prevented with the compositions of the inventtion include, but are not limited to, aortic valve stenosis, atrial septal defects, artioventricular (A-V) canal defect, bicuspid aortic valve, coarctation or the aorta, dextrocardia, Ebstein's anomaly, Eisenmenger's complex, hypoplastic left heart syndrome, Marfan syndrome, patent ductus arteriosus, progeria, pulmonary atresia, pulmonary valve stenosis, subaortic stenosis, tetralogy of fallot, total anomalous pulmonary venous (P-V) connection, transposition of the great arteries, tricuspid

atresia, truncus arteriosus, ventricular septal defects. Developmental or inherited disorders resulting in disorders involving the hematopoietic system that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but not limited to, Bernard-Soulier syndrome, Chédiak-Higashi syndrome, hemophilia, Hermansky-Pudlak syndrome, sickle cell anemia, storage pool disease, thromboxane A2 dysfunction, thrombasthenia, and von Willebrand's disease.

[0733] The compositions of the invention may also be used to diagnose, treat, and/or prevent developmental and inherited disorders resulting in disorders or defects of the gastrointestinal system, including, but not limited to, anal atresia, biliary atresia, esophageal atresia, diaphragmatic hernia, Hirschsprung's disease, Meckel's diverticulum, oligohydramnios, omphalocele, polyhydramnios, porphyria, situs inversus viscera. Developmental or inherited disorders resulting in metabolic disorders that may be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, alpha-1 antitrypsin deficiency, cystic fibrosis, hemochromatosis, lysosomal storage disease, phenylketonuria, Wilson's disease, and Zellweger syndrome.

[0734] Disorders of the reproductive system that are developmentally or hereditary related that may also be diagnosed, treated, and/or prevented with the compositions of the invention include, but are not limited to, androgen insensitivity syndrome, ambiguous genitalia, autosomal sex reversal, congenital adreneal hyperplasia, gonadoblastoma, ovarian germ cell cancer. pseudohermphroditism. hermaphroditism, undescended testis, XX male syndrome, and XY female type gonadal dysgenesis. The compositions of the invention may also be used to diagnose, treat, and/or prevent developmental or inherited respiratory defects including, but not limited to, askin tumor, azygos lobe, congenital diaphragmatic hernia, congenital lobar emphysema, cystic adenomatoid malformation, lobar emphysema, hyaline membrane disease, and pectus excavatum.

[0735] Developmental or inherited disorders may also result from chromosomal or genetic aberration that may be diagnosed, treated, and/or prevented with the compositions of the invention including, but not limited to, 4p- syndrome, cri du chat syndrome, Digeorge syndrome, Down's syndrome, Edward's syndrome, fragile X syndrome, Klinefelter's syndrome, Patau's syndrome, Prader-Willi syndrome,

progeria, Turner's syndrome, triple X syndrome, and XYY syndrome. Other developmental disorders that can be diagnosed, treated, and/or prevented with the compositions of the invention, include, but are not limited to, fetal alcohol syndrome, and can be caused by environmental factors surrounding the developing fetus.

[0736] The compositions of the invention may further be able to be used to diagnose, treat, and/or prevent errors in development or a genetic disposition that may result in hyperproliferative disorders or neoplasms, including, but not limited to, acute childhood lymphoblastic leukemia, askin tumor, Beckwith-Wiedemann syndrome, childhood acute myeloid leukemia, childhood brain stem glioma, childhood cerebellar astrocytoma, childhood extracranial germ cell tumors childhood (primary), gonadoblastoma, hepatocellular cancer, childhood Hodgkin's disease, childhood Hodgkin's lymphoma, childhood hypothalamic and visual pathway glioma, childhood (primary) liver cancer, childhood lymphoblastic leukemia. childhood medulloblastoma, childhood non-Hodgkin's lymphoma, childhood pineal and supratentorial primitive neuroectodermal tumors, childhood primary liver cancer, childhood rhabdomyosarcoma, childhood soft tissue sarcoma, Gorlin syndrome, familial multiple endrocrine neoplasia type I, neuroblastoma, ovarian germ cell cancer, pheochromocytoma, retinoblastoma, and Wilm's tumor.

[0737] Polypeptides may be administered using any method known in the art, including, but not limited to, direct needle injection at the delivery site, intravenous injection, topical administration, catheter infusion, biolistic injectors, particle accelerators, gelfoam sponge depots, other commercially available depot materials, osmotic pumps, oral or suppositorial solid pharmaceutical formulations, decanting or topical applications during surgery, aerosol delivery. Such methods are known in the art. Polypeptides may be administered as part of a Therapeutic, described in more detail below. Methods of delivering polynucleotides are described in more detail herein.

Diseases at the Cellular Level

[0738] Diseases associated with increased cell survival or the inhibition of apoptosis that could be treated, prevented, diagnosed and/or prognosed using polynucleotides or polypeptides, as well as antagonists or agonists of the present

invention, include cancers (such as follicular lymphomas, carcinomas with p53 mutations, and hormone-dependent tumors, including, but not limited to colon cancer, cardiac tumors, pancreatic cancer, melanoma, retinoblastoma, glioblastoma, lung cancer, intestinal cancer, testicular cancer, stomach cancer, neuroblastoma, myxoma, myoma, lymphoma, endothelioma, osteoblastoma, osteoclastoma, osteosarcoma, chondrosarcoma, adenoma, breast cancer, prostate cancer, Kaposi's sarcoma and ovarian cancer); autoimmune disorders (such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis) and viral infections (such as herpes viruses, pox viruses and adenoviruses), inflammation, graft v. host disease, acute graft rejection, and chronic graft rejection.

[0739] In preferred embodiments, polynucleotides, polypeptides, and/or antagonists of the invention are used to inhibit growth, progression, and/or metastasis of cancers, in particular those [listed above] involving reproductive system tissues.

Additional diseases or conditions associated with increased cell survival [0740] that could be treated or detected by polynucleotides or polypeptides, or agonists or antagonists of the present invention include, but are not limited to, progression, and/or metastases of malignancies and related disorders such as leukemia (including acute leukemias (e.g., acute lymphocytic leukemia, acute myelocytic leukemia (including myeloblastic, promyelocytic, myelomonocytic, monocytic, and erythroleukemia)) and chronic leukemias (e.g., chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia)), polycythemia vera, lymphomas (e.g., Hodgkin's disease and non-Hodgkin's disease), multiple myeloma, Waldenstrom's macroglobulinemia, heavy chain disease, and solid tumors including, but not limited to, sarcomas and carcinomas such as fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, chordoma. angiosarcoma, endotheliosarcoma, lymphangiosarcoma, lymphangioendotheliosarcoma, synovioma, mesothelioma. Ewing's tumor, leiomyosarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary adenocarcinomas, cystadenocarcinoma, medullary

carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma, choriocarcinoma, seminoma, embryonal carcinoma, Wilm's tumor, cervical cancer, testicular tumor, lung carcinoma, small cell lung carcinoma, bladder carcinoma, epithelial carcinoma, glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, pinealoma, hemangioblastoma, acoustic neuroma, oligodendroglioma, menangioma, melanoma, neuroblastoma, and retinoblastoma.

prevented, diagnosted, and/or prognosed using polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, include, but are not limited to, AIDS; neurodegenerative disorders (such as Alzheimer's disease, Parkinson's disease, Amyotrophic lateral sclerosis, Retinitis pigmentosa, Cerebellar degeneration and brain tumor or prior associated disease); autoimmune disorders (such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis) myelodysplastic syndromes (such as aplastic anemia), graft v. host disease, ischemic injury (such as that caused by myocardial infarction, stroke and reperfusion injury), liver injury (e.g., hepatitis related liver injury, ischemia/reperfusion injury, cholestosis (bile duct injury) and liver cancer); toxin-induced liver disease (such as that caused by alcohol), septic shock, cachexia and anorexia.

Wound Healing and Epithelial Cell Proliferation

In accordance with yet a further aspect of the present invention, there is provided a process for utilizing polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, for therapeutic purposes, for example, to stimulate epithelial cell proliferation and basal keratinocytes for the purpose of wound healing, and to stimulate hair follicle production and healing of dermal wounds. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, may be clinically useful in stimulating wound healing including surgical wounds, excisional wounds, deep wounds involving damage of the dermis and epidermis, eye tissue wounds, dental tissue wounds, oral cavity wounds, diabetic ulcers, dermal ulcers, cubitus ulcers, arterial ulcers, venous stasis ulcers, burns

resulting from heat exposure or chemicals, and other abnormal wound healing conditions such as uremia, malnutrition, vitamin deficiencies and complications associated with systemic treatment with steroids, radiation therapy and antineoplastic drugs and antimetabolites. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to promote dermal reestablishment subsequent to dermal loss.

Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to increase the adherence of skin grafts to a wound bed and to stimulate re-epithelialization from the wound bed. The following are types of grafts that polynucleotides or polypeptides, agonists or antagonists of the present invention, could be used to increase adherence to a wound bed: autografts, artificial skin, allografts, autodermic graft, autoepdermic grafts, avacular grafts, Blair-Brown grafts, bone graft, brephoplastic grafts, cutis graft, delayed graft, dermic graft, epidermic graft, fascia graft, full thickness graft, heterologous graft, xenograft, homologous graft, hyperplastic graft, lamellar graft, mesh graft, mucosal graft, Ollier-Thiersch graft, omenpal graft, patch graft, pedicle graft, penetrating graft, split skin graft, thick split graft. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, can be used to promote skin strength and to improve the appearance of aged skin.

It is believed that polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, will also produce changes in hepatocyte proliferation, and epithelial cell proliferation in the lung, breast, pancreas, stomach, small intestine, and large intestine. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could promote proliferation of epithelial cells such as sebocytes, hair follicles, hepatocytes, type II pneumocytes, mucin-producing goblet cells, and other epithelial cells and their progenitors contained within the skin, lung, liver, and gastrointestinal tract. Polynucleotides or polypeptides, agonists or antagonists of the present invention, may promote proliferation of endothelial cells, keratinocytes, and basal keratinocytes.

[0745] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could also be used to reduce the side effects of gut toxicity that result from radiation, chemotherapy treatments or viral infections. Polynucleotides or

polypeptides, as well as agonists or antagonists of the present invention, may have a cytoprotective effect on the small intestine mucosa. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, may also stimulate healing of mucositis (mouth ulcers) that result from chemotherapy and viral infections.

[0746] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could further be used in full regeneration of skin in full and partial thickness skin defects, including burns, (i.e., repopulation of hair follicles, sweat glands, and sebaceous glands), treatment of other skin defects such as psoriasis. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to treat epidermolysis bullosa, a defect in adherence of the epidermis to the underlying dermis which results in frequent, open and painful blisters by accelerating reepithelialization of these lesions. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could also be used to treat gastric and doudenal ulcers and help heal by scar formation of the mucosal lining and regeneration of glandular mucosa and duodenal mucosal lining more rapidly. Inflammatory bowel diseases, such as Crohn's disease and ulcerative colitis, are diseases, which result in destruction of the mucosal surface of the small or large intestine, respectively. Thus, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to promote the resurfacing of the mucosal surface to aid more rapid healing and to prevent progression of inflammatory bowel disease. Treatment with polynucleotides or polypeptides, agonists or antagonists of the present invention, is expected to have a significant effect on the production of mucus throughout the gastrointestinal tract and could be used to protect the intestinal mucosa from injurious substances that are ingested or following surgery. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to treat diseases associate with the under expression.

[0747] Moreover, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to prevent and heal damage to the lungs due to various pathological states. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could stimulate proliferation and differentiation and promote the repair of alveoli and brochiolar epithelium to prevent or treat acute or chronic lung damage. For example, emphysema, which results in the

progressive loss of aveoli, and inhalation injuries, i.e., resulting from smoke inhalation and burns, that cause necrosis of the bronchiolar epithelium and alveoli could be effectively treated using polynucleotides or polypeptides, agonists or antagonists of the present invention. Also, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to stimulate the proliferation of and differentiation of type II pneumocytes, which may help treat or prevent disease such as hyaline membrane diseases, such as infant respiratory distress syndrome and bronchopulmonary displasia, in premature infants.

[0748] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could stimulate the proliferation and differentiation of hepatocytes and, thus, could be used to alleviate or treat liver diseases and pathologies such as fulminant liver failure caused by cirrhosis, liver damage caused by viral hepatitis and toxic substances (i.e., acetaminophen, carbon tetraholoride and other hepatotoxins known in the art).

In addition, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used treat or prevent the onset of diabetes mellitus. In patients with newly diagnosed Types I and II diabetes, where some islet cell function remains, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to maintain the islet function so as to alleviate, delay or prevent permanent manifestation of the disease. Also, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used as an auxiliary in islet cell transplantation to improve or promote islet cell function.

Infectious Diseases

[0749] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention can be used to treat or detect infectious agents. For example, by increasing the immune response, particularly increasing the proliferation and differentiation of B and/or T cells, infectious diseases may be treated. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may also directly inhibit the infectious agent, without necessarily eliciting an immune response.

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Viruses are one example of an infectious agent that can cause disease or [0750] symptoms that can be treated or detected by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention. Examples of viruses, include, but are not limited to Examples of viruses, include, but are not limited to the following DNA and RNA viruses and viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus, Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Dengue, EBV, HIV, Flaviviridae, Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus, Rhabdoviridae), Orthomyxoviridae (e.g., Influenza A, Influenza B, and parainfluenza), Papiloma virus, Papovaviridae, Parvoviridae, Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g., Rotavirus), Retroviridae (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). Viruses falling within these families can cause a variety of diseases or symptoms, including, but not limited to: arthritis, bronchiollitis, respiratory syncytial virus, encephalitis, eye infections (e.g., conjunctivitis, keratitis), chronic fatigue syndrome, hepatitis (A, B, C, E, Chronic Active, Delta), Japanese B encephalitis, Junin, Chikungunya, Rift Valley fever, yellow fever, meningitis, opportunistic infections (e.g., AIDS), pneumonia, Burkitt's Lymphoma, chickenpox, hemorrhagic fever, Measles, Mumps, Parainfluenza, Rabies, the common cold, Polio, leukemia, Rubella, sexually transmitted diseases, skin diseases (e.g., Kaposi's, warts), and viremia. polynucleotides or polypeptides, or agonists or antagonists of the invention, can be used to treat or detect any of these symptoms or diseases. In specific embodiments, polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat: meningitis, Dengue, EBV, and/or hepatitis (e.g., hepatitis B). In an additional specific embodiment polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat patients nonresponsive to one or more other commercially available hepatitis vaccines. In a further specific embodiment polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat AIDS.

[0751] Similarly, bacterial and fungal agents that can cause disease or symptoms and that can be treated or detected by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention include, but not limited to, the following Gram-

Negative and Gram-positive bacteria, bacterial families, and fungi: Actinomyces (e.g., Norcardia), Acinetobacter, Cryptococcus neoformans, Aspergillus, Bacillaceae (e.g., Bacillus anthrasis), Bacteroides (e.g., Bacteroides fragilis), Blastomycosis, Bordetella, Borrelia (e.g., Borrelia burgdorferi), Brucella, Candidia, Campylobacter, Chlamydia, Clostridium (e.g., Clostridium botulinum, Clostridium dificile, Clostridium perfringens, Clostridium tetani), Coccidioides, Corynebacterium (e.g., Corynebacterium diptheriae), Cryptococcus, Dermatocycoses, E. coli (e.g., Enterotoxigenic E. coli and Enterohemorrhagic E. coli), Enterobacter (e.g. Enterobacter aerogenes), Enterobacteriaceae (Klebsiella, Salmonella (e.g., Salmonella typhi, Salmonella enteritidis, Salmonella typhi), Serratia, Yersinia, Shigella), Erysipelothrix, Haemophilus (e.g., Haemophilus influenza type B), Helicobacter, Legionella (e.g., Legionella pneumophila), Leptospira, Listeria (e.g., Listeria monocytogenes), Mycoplasma, Mycobacterium (e.g., Mycobacterium leprae and Mycobacterium tuberculosis), Vibrio (e.g., Vibrio cholerae), Neisseriaceae (e.g., Neisseria gonorrhea, Neisseria meningitidis), Pasteurellacea, Proteus, Pseudomonas (e.g., Pseudomonas aeruginosa), Rickettsiaceae, Spirochetes (e.g., Treponema spp., Leptospira spp., Borrelia spp.), Shigella spp., Staphylococcus (e.g., Staphylococcus aureus), Meningiococcus, Pneumococcus and Streptococcus (e.g., Streptococcus pneumoniae and Groups A, B, and C Streptococci), and Ureaplasmas. These bacterial, parasitic, and fungal families can cause diseases or symptoms, including, but not limited to: antibiotic-resistant infections, bacteremia, endocarditis, septicemia, eye infections (e.g., conjunctivitis), uveitis, tuberculosis, gingivitis, bacterial diarrhea, opportunistic infections (e.g., AIDS related infections), paronychia, prosthesis-related infections, dental caries, Reiter's Disease, respiratory tract infections, such as Whooping Cough or Empyema, sepsis, Lyme Disease, Cat-Scratch Disease, dysentery, paratyphoid fever, food poisoning, Legionella disease, chronic and acute inflammation, erythema, yeast infections, typhoid, pneumonia, gonorrhea, meningitis (e.g., mengitis types A and B), chlamydia, syphillis, diphtheria, leprosy, brucellosis, peptic ulcers, anthrax, spontaneous abortions, birth defects, pneumonia, lung infections, ear infections, deafness, blindness, lethargy, malaise, vomiting, chronic diarrhea, Crohn's disease, colitis, vaginosis, sterility, pelvic inflammatory diseases, candidiasis, paratuberculosis, tuberculosis, lupus, botulism, gangrene, tetanus,

impetigo, Rheumatic Fever, Scarlet Fever, sexually transmitted diseases, skin diseases (e.g., cellulitis, dermatocycoses), toxemia, urinary tract infections, wound infections, noscomial infections. Polynucleotides or polypeptides, agonists or antagonists of the invention, can be used to treat or detect any of these symptoms or diseases. In specific embodiments, polynucleotides, polypeptides, agonists or antagonists of the invention are used to treat: tetanus, diptheria, botulism, and/or meningitis type B.

Moreover, parasitic agents causing disease or symptoms that can be treated, [0752] prevented, and/or diagnosed by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention include, but not limited to, the following families or class: Amebiasis, Babesiosis, Coccidiosis, Cryptosporidiosis, Dientamoebiasis, Ectoparasitic, Giardias, Helminthiasis, Leishmaniasis, Dourine, Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas and Sporozoans (e.g., Plasmodium virax, Plasmodium falciparium, Plasmodium malariae and Plasmodium ovale). These parasites can cause a variety of diseases or symptoms, including, but not limited to: Scabies, Trombiculiasis, eye infections, intestinal disease (e.g., dysentery, giardiasis), liver disease, lung disease, opportunistic infections (e.g., AIDS related), malaria, pregnancy complications, and toxoplasmosis. polynucleotides or polypeptides, or agonists or antagonists of the invention, can be used to treat, prevent, and/or diagnose any of these symptoms or diseases. In specific embodiments, polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat, prevent, and/or diagnose malaria.

[0753] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention of the present invention could either be by administering an effective amount of a polypeptide to the patient, or by removing cells from the patient, supplying the cells with a polynucleotide of the present invention, and returning the engineered cells to the patient (ex vivo therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen in a vaccine to raise an immune response against infectious disease.

Regeneration

[0754] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention can be used to differentiate, proliferate, and attract cells, leading to

the regeneration of tissues. (See, Science 276:59-87 (1997).) The regeneration of tissues could be used to repair, replace, or protect tissue damaged by congenital defects, trauma (wounds, burns, incisions, or ulcers), age, disease (e.g. osteoporosis, osteocarthritis, periodontal disease, liver failure), surgery, including cosmetic plastic surgery, fibrosis, reperfusion injury, or systemic cytokine damage.

organs (e.g., pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac), vasculature (including vascular and lymphatics), nervous, hematopoietic, and skeletal (bone, cartilage, tendon, and ligament) tissue. Preferably, regeneration occurs without or decreased scarring. Regeneration also may include angiogenesis.

Moreover, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, may increase regeneration of tissues difficult to heal. For example, increased tendon/ligament regeneration would quicken recovery time after damage. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention could also be used prophylactically in an effort to avoid damage. Specific diseases that could be treated include of tendinitis, carpal tunnel syndrome, and other tendon or ligament defects. A further example of tissue regeneration of non-healing wounds includes pressure ulcers, ulcers associated with vascular insufficiency, surgical, and traumatic wounds.

[0757] Similarly, nerve and brain tissue could also be regenerated by using polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, to proliferate and differentiate nerve cells. Diseases that could be treated using this method include central and peripheral nervous system diseases, neuropathies, or mechanical and traumatic disorders (e.g., spinal cord disorders, head trauma, cerebrovascular disease, and stoke). Specifically, diseases associated with peripheral nerve injuries, peripheral neuropathy (e.g., resulting from chemotherapy or other medical therapies), localized neuropathies, and central nervous system diseases (e.g., Alzheimer's disease, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome), could all be treated using the polynucleotides or polypeptides, as well as agonists or antagonists of the present invention.

Chemotaxis

[0758] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may have chemotaxis activity. A chemotaxic molecule attracts or mobilizes cells (e.g., monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells) to a particular site in the body, such as inflammation, infection, or site of hyperproliferation. The mobilized cells can then fight off and/or heal the particular trauma or abnormality.

[0759] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may increase chemotaxic activity of particular cells. These chemotactic molecules can then be used to treat inflammation, infection, hyperproliferative disorders, or any immune system disorder by increasing the number of cells targeted to a particular location in the body. For example, chemotaxic molecules can be used to treat wounds and other trauma to tissues by attracting immune cells to the injured location. Chemotactic molecules of the present invention can also attract fibroblasts, which can be used to treat wounds.

[0760] It is also contemplated that polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may inhibit chemotactic activity. These molecules could also be used to treat disorders. Thus, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention could be used as an inhibitor of chemotaxis.

Binding Activity

[0761] A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or small molecules.

[0762] Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural or functional mimetic. (See, Coligan et al., Current Protocols in Immunology

1(2):Chapter 5 (1991).) Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

[0763] Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide. Preferred cells include cells from mammals, yeast, Drosophila, or *E. coli*. Cells expressing the polypeptide (or cell membrane containing the expressed polypeptide) are then preferably contacted with a test compound potentially containing the molecule to observe binding, stimulation, or inhibition of activity of either the polypeptide or the molecule.

[0764] The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the candidate compound results in a signal generated by binding to the polypeptide.

[0765] Alternatively, the assay can be carried out using cell-free preparations, polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule activity or binding, and comparing the polypeptide/molecule activity or binding to a standard.

[0766] Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

[0767] Additionally, the receptor to which the polypeptide of the present invention binds can be identified by numerous methods known to those of skill in the art, for example, ligand panning and FACS sorting (Coligan, et al., Current Protocols in Immun., 1(2), Chapter 5, (1991)). For example, expression cloning is employed wherein polyadenylated RNA is prepared from a cell responsive to the polypeptides, for example, NIH3T3 cells which are known to contain multiple receptors for the FGF family proteins, and SC-3 cells, and a cDNA library created from this RNA is divided into pools and used to transfect COS cells or other cells that are not responsive to the

polypeptides. Transfected cells which are grown on glass slides are exposed to the polypeptide of the present invention, after they have been labeled. The polypeptides can be labeled by a variety of means including iodination or inclusion of a recognition site for a site-specific protein kinase.

[0768] Following fixation and incubation, the slides are subjected to autoradiographic analysis. Positive pools are identified and sub-pools are prepared and retransfected using an iterative sub-pooling and re-screening process, eventually yielding a single clones that encodes the putative receptor.

[0769] As an alternative approach for receptor identification, the labeled polypeptides can be photoaffinity linked with cell membrane or extract preparations that express the receptor molecule. Cross-linked material is resolved by PAGE analysis and exposed to X-ray film. The labeled complex containing the receptors of the polypeptides can be excised, resolved into peptide fragments, and subjected to protein microsequencing. The amino acid sequence obtained from microsequencing would be used to design a set of degenerate oligonucleotide probes to screen a cDNA library to identify the genes encoding the putative receptors.

Moreover, the techniques of gene-shuffling, motif-shuffling, exon-[0770] shuffling, and/or codon-shuffling (collectively referred to as "DNA shuffling") may be employed to modulate the activities of the polypeptide of the present invention thereby effectively generating agonists and antagonists of the polypeptide of the present See generally, U.S. Patent Nos. 5,605,793, 5,811,238, 5,830,721, invention. 5,834,252, and 5,837,458, and Patten, P. A., et al., Curr. Opinion Biotechnol. 8:724-33 (1997); Harayama, S. Trends Biotechnol. 16(2):76-82 (1998); Hansson L. O., et al., J. Mol. Biol. 287:265-76 (1999); and Lorenzo, M. M. and Blasco, R. Biotechniques 24(2):308-13 (1998); each of these patents and publications are hereby incorporated by reference). In one embodiment, alteration of polynucleotides and corresponding polypeptides may be achieved by DNA shuffling. DNA shuffling involves the assembly of two or more DNA segments into a desired molecule by homologous, or site-specific, recombination. In another embodiment, polynucleotides and corresponding polypeptides may be altered by being subjected to random mutagenesis by error-prone PCR, random nucleotide insertion or other methods prior to recombination. In another embodiment, one or more components, motifs, sections,

parts, domains, fragments, etc., of the polypeptide of the present invention may be recombined with one or more components, motifs, sections, parts, domains, fragments, etc. of one or more heterologous molecules. In preferred embodiments, the heterologous molecules are family members. In further preferred embodiments, the heterologous molecule is a growth factor such as, for example, platelet-derived growth factor (PDGF), insulin-like growth factor (IGF-I), transforming growth factor (TGF)-alpha, epidermal growth factor (EGF), fibroblast growth factor (FGF), TGF-beta, bone morphogenetic protein (BMP)-2, BMP-4, BMP-5, BMP-6, BMP-7, activins A and B, decapentaplegic(dpp), 60A, OP-2, dorsalin, growth differentiation factors (GDIFs), nodal, MIS, inhibin-alpha, TGF-beta1, TGF-beta2, TGF-beta3, TGF-beta5, and glial-derived neurotrophic factor (GDNF).

[0771] Other preferred fragments are biologically active fragments of the polypeptide of the present invention. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

identify those which modulate the action of the polypeptide of the present invention. An example of such an assay comprises combining a mammalian fibroblast cell, the polypeptide of the present invention, the compound to be screened and ³[H] thymidine under cell culture conditions where the fibroblast cell would normally proliferate. A control assay may be performed in the absence of the compound to be screened and compared to the amount of fibroblast proliferation in the presence of the compound to determine if the compound stimulates proliferation by determining the uptake of ³[H] thymidine in each case. The amount of fibroblast cell proliferation is measured by liquid scintillation chromatography which measures the incorporation of ³[H] thymidine. Both agonist and antagonist compounds may be identified by this procedure.

[0773] In another method, a mammalian cell or membrane preparation expressing a receptor for a polypeptide of the present invention is incubated with a labeled polypeptide of the present invention in the presence of the compound. The ability of

the compound to enhance or block this interaction could then be measured. Alternatively, the response of a known second messenger system following interaction of a compound to be screened and the receptor is measured and the ability of the compound to bind to the receptor and elicit a second messenger response is measured to determine if the compound is a potential agonist or antagonist. Such second messenger systems include but are not limited to, cAMP guanylate cyclase, ion channels or phosphoinositide hydrolysis.

[0774] All of these above assays can be used as diagnostic or prognostic markers. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptides of the invention from suitably manipulated cells or tissues.

[0775] Therefore, the invention includes a method of identifying compounds which bind to a polypeptide of the invention comprising the steps of: (a) incubating a candidate binding compound with a polypeptide of the present invention; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound with a polypeptide of the present invention, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

Targeted Delivery

[0776] In another embodiment, the invention provides a method of delivering compositions to targeted cells expressing a receptor for a polypeptide of the invention, or cells expressing a cell bound form of a polypeptide of the invention.

[0777] As discussed herein, polypeptides or antibodies of the invention may be associated with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs via hydrophobic, hydrophilic, ionic and/or covalent interactions. In one embodiment, the invention provides a method for the specific delivery of compositions of the invention to cells by administering polypeptides of the invention (including antibodies) that are associated with heterologous polypeptides or nucleic acids. In one

example, the invention provides a method for delivering a therapeutic protein into the targeted cell. In another example, the invention provides a method for delivering a single stranded nucleic acid (e.g., antisense or ribozymes) or double stranded nucleic acid (e.g., DNA that can integrate into the cell's genome or replicate episomally and that can be transcribed) into the targeted cell.

[0778] In another embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention (e.g., polypeptides of the invention or antibodies of the invention) in association with toxins or cytotoxic prodrugs.

By "toxin" is meant compounds that bind and activate endogenous [0779]cytotoxic effector systems, radioisotopes, holotoxins, modified toxins, catalytic subunits of toxins, or any molecules or enzymes not normally present in or on the surface of a cell that under defined conditions cause the cell's death. Toxins that may be used according to the methods of the invention include, but are not limited to, radioisotopes known in the art, compounds such as, for example, antibodies (or complement fixing containing portions thereof) that bind an inherent or induced endogenous cytotoxic effector system, thymidine kinase, endonuclease, RNAse, alpha toxin, ricin, abrin, Pseudomonas exotoxin A, diphtheria toxin, saporin, momordin, gelonin, pokeweed antiviral protein, alpha-sarcin and cholera toxin. By "cytotoxic prodrug" is meant a non-toxic compound that is converted by an enzyme, normally present in the cell, into a cytotoxic compound. Cytotoxic prodrugs that may be used according to the methods of the invention include, but are not limited to, glutamyl derivatives of benzoic acid mustard alkylating agent, phosphate derivatives of etoposide or mitomycin C, cytosine arabinoside, daunorubisin, and phenoxyacetamide derivatives of doxorubicin.

Drug Screening

[0780] Further contemplated is the use of the polypeptides of the present invention, or the polynucleotides encoding these polypeptides, to screen for molecules which modify the activities of the polypeptides of the present invention. Such a method would include contacting the polypeptide of the present invention with a selected compound(s) suspected of having antagonist or agonist activity, and assaying

the activity of these polypeptides following binding.

[0781] This invention is particularly useful for screening therapeutic compounds by using the polypeptides of the present invention, or binding fragments thereof, in any of a variety of drug screening techniques. The polypeptide or fragment employed in such a test may be affixed to a solid support, expressed on a cell surface, free in solution, or located intracellularly. One method of drug screening utilizes eukaryotic or prokaryotic host cells which are stably transformed with recombinant nucleic acids expressing the polypeptide or fragment. Drugs are screened against such transformed cells in competitive binding assays. One may measure, for example, the formulation of complexes between the agent being tested and a polypeptide of the present invention.

Other agents which affect activities mediated by the polypeptides of the present invention. These methods comprise contacting such an agent with a polypeptide of the present invention or a fragment thereof and assaying for the presence of a complex between the agent and the polypeptide or a fragment thereof, by methods well known in the art. In such a competitive binding assay, the agents to screen are typically labeled. Following incubation, free agent is separated from that present in bound form, and the amount of free or uncomplexed label is a measure of the ability of a particular agent to bind to the polypeptides of the present invention.

[0783] Another technique for drug screening provides high throughput screening for compounds having suitable binding affinity to the polypeptides of the present invention, and is described in great detail in European Patent Application 84/03564, published on September 13, 1984, which is incorporated herein by reference herein. Briefly stated, large numbers of different small peptide test compounds are synthesized on a solid substrate, such as plastic pins or some other surface. The peptide test compounds are reacted with polypeptides of the present invention and washed. Bound polypeptides are then detected by methods well known in the art. Purified polypeptides are coated directly onto plates for use in the aforementioned drug screening techniques. In addition, non-neutralizing antibodies may be used to capture the peptide and immobilize it on the solid support.

[0784] This invention also contemplates the use of competitive drug screening assays in which neutralizing antibodies capable of binding polypeptides of the present

invention specifically compete with a test compound for binding to the polypeptides or fragments thereof. In this manner, the antibodies are used to detect the presence of any peptide which shares one or more antigenic epitopes with a polypeptide of the invention.

Antisense And Ribozyme (Antagonists)

In specific embodiments, antagonists according to the present invention are [0785] nucleic acids corresponding to the sequences contained in SEQ ID NO:X, or the complementary strand thereof, and/or to cDNA sequences contained in cDNA Clone ID NO:Z identified for example, in Table 1A. In one embodiment, antisense sequence is generated internally, by the organism, in another embodiment, the antisense sequence is separately administered (see, for example, O'Connor, J., Neurochem. 56:560 (1991). Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988). Antisense technology can be used to control gene expression through antisense DNA or RNA, or through 'triple-helix formation. Antisense techniques are discussed for example, in Okano, J., Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988). Triple helix formation is discussed in, for instance, Lee et al., Nucleic Acids Research 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1300 (1991). The methods are based on binding of a polynucleotide to a complementary DNA or RNA.

[0786] For example, the use of c-myc and c-myb antisense RNA constructs to inhibit the growth of the non-lymphocytic leukemia cell line HL-60 and other cell lines was previously described. (Wickstrom et al. (1988); Anfossi et al. (1989)). These experiments were performed in vitro by incubating cells with the oligoribonucleotide. A similar procedure for in vivo use is described in WO 91/15580. Briefly, a pair of oligonucleotides for a given antisense RNA is produced as follows: A sequence complimentary to the first 15 bases of the open reading frame is flanked by an EcoR1 site on the 5' end and a HindIII site on the 3' end. Next, the pair of oligonucleotides is heated at 90°C for one minute and then annealed in 2X ligation buffer (20mM TRIS HCl pH 7.5, 10mM MgCl2, 10MM dithiothreitol (DTT)

and 0.2 mM ATP) and then ligated to the EcoR1/Hind III site of the retroviral vector PMV7 (WO 91/15580).

[0787] For example, the 5' coding portion of a polynucleotide that encodes the polypeptide of the present invention may be used to design an antisense RNA oligonucleotide of from about 10 to 40 base pairs in length. A DNA oligonucleotide is designed to be complementary to a region of the gene involved in transcription thereby preventing transcription and the production of the receptor. The antisense RNA oligonucleotide hybridizes to the mRNA in vivo and blocks translation of the mRNA molecule into receptor polypeptide.

[0788] In one embodiment, the antisense nucleic acid of the invention is produced intracellularly by transcription from an exogenous sequence. For example, a vector or a portion thereof, is transcribed, producing an antisense nucleic acid (RNA) of the invention. Such a vector would contain a sequence encoding the antisense nucleic acid. Such a vector can remain episomal or become chromosomally integrated, as long as it can be transcribed to produce the desired antisense RNA. Such vectors can be constructed by recombinant DNA technology methods standard in the art. Vectors can be plasmid, viral, or others known in the art, used for replication and expression in vertebrate cells. Expression of the sequence encoding the polypeptide of the present invention or fragments thereof, can be by any promoter known in the art to act in vertebrate, preferably human cells. Such promoters can be inducible or constitutive. Such promoters include, but are not limited to, the SV40 early promoter region (Bernoist and Chambon, Nature 29:304-310 (1981), the promoter contained in the 3' long terminal repeat of Rous sarcoma virus (Yamamoto et al., Cell 22:787-797 (1980), the herpes thymidine promoter (Wagner et al., Proc. Natl. Acad. Sci. U.S.A. 78:1441-1445 (1981), the regulatory sequences of the metallothionein gene (Brinster, et al., Nature 296:39-42 (1982)), etc.

[0789] The antisense nucleic acids of the invention comprise a sequence complementary to at least a portion of an RNA transcript of a gene of the present invention. However, absolute complementarity, although preferred, is not required. A sequence "complementary to at least a portion of an RNA," referred to herein, means a sequence having sufficient complementarity to be able to hybridize with the RNA, forming a stable duplex; in the case of double stranded antisense nucleic acids, a single

strand of the duplex DNA may thus be tested, or triplex formation may be assayed. The ability to hybridize will depend on both the degree of complementarity and the length of the antisense nucleic acid. Generally, the larger the hybridizing nucleic acid, the more base mismatches with a RNA it may contain and still form a stable duplex (or triplex as the case may be). One skilled in the art can ascertain a tolerable degree of mismatch by use of standard procedures to determine the melting point of the hybridized complex.

Oligonucleotides that are complementary to the 5' end of the message, e.g., [0790] the 5' untranslated sequence up to and including the AUG initiation codon, should work most efficiently at inhibiting translation. However, sequences complementary to the 3' untranslated sequences of mRNAs have been shown to be effective at inhibiting translation of mRNAs as well. See generally, Wagner, R., 1994, Nature 372:333-335. Thus, oligonucleotides complementary to either the 5'- or 3'- non- translated, noncoding regions of polynucleotide sequences described herein could be used in an antisense approach to inhibit translation of endogenous mRNA. Oligonucleotides complementary to the 5' untranslated region of the mRNA should include the complement of the AUG start codon. Antisense oligonucleotides complementary to mRNA coding regions are less efficient inhibitors of translation but could be used in accordance with the invention. Whether designed to hybridize to the 5'-, 3'- or coding region of mRNA of the present invention, antisense nucleic acids should be at least six nucleotides in length, and are preferably oligonucleotides ranging from 6 to about 50 nucleotides in length. In specific aspects the oligonucleotide is at least 10 nucleotides, at least 17 nucleotides, at least 25 nucleotides or at least 50 nucleotides.

[0791] The polynucleotides of the invention can be DNA or RNA or chimeric mixtures or derivatives or modified versions thereof, single-stranded or double-stranded. The oligonucleotide can be modified at the base moiety, sugar moiety, or phosphate backbone, for example, to improve stability of the molecule, hybridization, etc. The oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors *in vivo*), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86:6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci. 84:648-652; PCT Publication No. WO88/09810, published December 15, 1988) or the blood-brain barrier (see, e.g., PCT

Publication No. WO89/10134, published April 25, 1988), hybridization-triggered cleavage agents. (See, e.g., Krol et al., 1988, BioTechniques 6:958-976) or intercalating agents. (See, e.g., Zon, 1988, Pharm. Res. 5:539-549). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, hybridization triggered cross-linking agent, transport agent, hybridization-triggered cleavage agent, etc.

The antisense oligonucleotide may comprise at least one modified base [0792] moiety which is selected from the group including, but not limited to, 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xantine, 4-acetylcytosine, 5-(carboxyhydroxylmethyl) uracil. 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine. 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine. 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil. 2-methylthio-N6isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine.

[0793] The antisense oligonucleotide may also comprise at least one modified sugar moiety selected from the group including, but not limited to, arabinose, 2-fluoroarabinose, xylulose, and hexose.

[0794] In yet another embodiment, the antisense oligonucleotide comprises at least one modified phosphate backbone selected from the group including, but not limited to, a phosphorothicate, a phosphorodithicate, a phosphoramidate, a phosphoramidate, a phosphoramidate, a methylphosphonate, an alkyl phosphotriester, and a formacetal or analog thereof.

[0795] In yet another embodiment, the antisense oligonucleotide is an a-anomeric oligonucleotide. An a-anomeric oligonucleotide forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual b-units, the strands run parallel to each other (Gautier et al., 1987, Nucl. Acids Res. 15:6625-6641). The

oligonucleotide is a 2'-0-methylribonucleotide (Inoue et al., 1987, Nucl. Acids Res. 15:6131-6148), or a chimeric RNA-DNA analogue (Inoue et al., 1987, FEBS Lett. 215:327-330).

[0796] Polynucleotides of the invention may be synthesized by standard methods known in the art, e.g. by use of an automated DNA synthesizer (such as are commercially available from Biosearch, Applied Biosystems, etc.). As examples, phosphorothioate oligonucleotides may be synthesized by the method of Stein et al. (1988, Nucl. Acids Res. 16:3209), methylphosphonate oligonucleotides can be prepared by use of controlled pore glass polymer supports (Sarin et al., 1988, Proc. Natl. Acad. Sci. U.S.A. 85:7448-7451), etc.

[0797] While antisense nucleotides complementary to the coding region sequence could be used, those complementary to the transcribed untranslated region are most preferred.

Potential antagonists according to the invention also include catalytic RNA, or a ribozyme (See, e.g., PCT International Publication WO 90/11364, published October 4, 1990; Sarver et al, Science 247:1222-1225 (1990). While ribozymes that cleave mRNA at site specific recognition sequences can be used to destroy mRNAs, the use of hammerhead ribozymes is preferred. Hammerhead ribozymes cleave mRNAs at locations dictated by flanking regions that form complementary base pairs with the target mRNA. The sole requirement is that the target mRNA have the following sequence of two bases: 5'-UG-3'. The construction and production of hammerhead ribozymes is well known in the art and is described more fully in Haseloff and Gerlach, Nature 334:585-591 (1988). There are numerous potential hammerhead ribozyme cleavage sites within the nucleotide sequence of SEQ ID NO:X. Preferably, the ribozyme is engineered so that the cleavage recognition site is located near the 5' end of the mRNA; i.e., to increase efficiency and minimize the intracellular accumulation of non-functional mRNA transcripts.

[0799] As in the antisense approach, the ribozymes of the invention can be composed of modified oligonucleotides (e.g. for improved stability, targeting, etc.) and should be delivered to cells which express in vivo. DNA constructs encoding the ribozyme may be introduced into the cell in the same manner as described above for the introduction of antisense encoding DNA. A preferred method of delivery involves

using a DNA construct "encoding" the ribozyme under the control of a strong constitutive promoter, such as, for example, pol III or pol II promoter, so that transfected cells will produce sufficient quantities of the ribozyme to destroy endogenous messages and inhibit translation. Since ribozymes unlike antisense molecules, are catalytic, a lower intracellular concentration is required for efficiency.

[0800] Antagonist/agonist compounds may be employed to inhibit the cell growth and proliferation effects of the polypeptides of the present invention on neoplastic cells and tissues, i.e. stimulation of angiogenesis of tumors, and, therefore, retard or prevent abnormal cellular growth and proliferation, for example, in tumor formation or growth.

[0801] The antagonist/agonist may also be employed to prevent hyper-vascular diseases, and prevent the proliferation of epithelial lens cells after extracapsular cataract surgery. Prevention of the mitogenic activity of the polypeptides of the present invention may also be desirous in cases such as restenosis after balloon angioplasty.

[0802] The antagonist/agonist may also be employed to prevent the growth of scar tissue during wound healing.

[0803] The antagonist/agonist may also be employed to treat the diseases described herein.

[0804] Thus, the invention provides a method of treating disorders or diseases, including but not limited to the disorders or diseases listed throughout this application, associated with overexpression of a polynucleotide of the present invention by administering to a patient (a) an antisense molecule directed to the polynucleotide of the present invention, and/or (b) a ribozyme directed to the polynucleotide of the present invention.

Binding Peptides and Other Molecules

[0805] The invention also encompasses screening methods for identifying polypeptides and nonpolypeptides that bind reproductive system related antigen polypeptides, and the reproductive system related antigen binding molecules identified thereby. These binding molecules are useful, for example, as agonists and antagonists of the reproductive system related antigen polypeptides. Such agonists and

antagonists can be used, in accordance with the invention, in the therapeutic embodiments described in detail, below.

[0806] This method comprises the steps of: contacting reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides with a plurality of molecules; and identifying a molecule that binds the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides.

The step of contacting the reproductive system related antigen polypeptides [0807] or reproductive system related antigen-like polypeptides with the plurality of molecules may be effected in a number of ways. For example, one may contemplate immobilizing the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides on a solid support and bringing a solution of the plurality of molecules in contact with the immobilized reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides. Such a procedure would be akin to an affinity chromatographic process, with the affinity matrix being comprised of the immobilized reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides. The molecules having a selective affinity for the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides can then be purified by affinity selection. The nature of the solid support, process for attachment of the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides to the solid support, solvent, and conditions of the affinity isolation or selection are largely conventional and well known to those of ordinary skill in the art.

[0808] Alternatively, one may also separate a plurality of polypeptides into substantially separate fractions comprising a subset of or individual polypeptides. For instance, one can separate the plurality of polypeptides by gel electrophoresis, column chromatography, or like method known to those of ordinary skill for the separation of polypeptides. The individual polypeptides can also be produced by a transformed host cell in such a way as to be expressed on or about its outer surface (e.g., a recombinant phage). Individual isolates can then be "probed" by the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides, optionally in the presence of an inducer should one be required for expression, to

determine if any selective affinity interaction takes place between the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides and the individual clone. Prior to contacting the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides with each fraction comprising individual polypeptides, the polypeptides could first be transferred to a solid support for additional convenience. Such a solid support may simply be a piece of filter membrane, such as one made of nitrocellulose or nylon. In this manner, positive clones could be identified from a collection of transformed host cells of an expression library, which harbor a DNA construct encoding a polypeptide having a selective affinity for reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides. Furthermore, the amino acid sequence of the polypeptide having a selective affinity for the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides can be determined directly by conventional means or the coding sequence of the DNA encoding the polypeptide can frequently be determined more conveniently. The primary sequence can then be deduced from the corresponding DNA sequence. If the amino acid sequence is to be determined from the polypeptide itself, one may use microsequencing techniques. The sequencing technique may include mass spectroscopy.

[0809] In certain situations, it may be desirable to wash away any unbound reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides, or alternatively, unbound polypeptides, from a mixture of the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides and the plurality of polypeptides prior to attempting to determine or to detect the presence of a selective affinity interaction. Such a wash step may be particularly desirable when the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides or the plurality of polypeptides is bound to a solid support.

[0810] The plurality of molecules provided according to this method may be provided by way of diversity libraries, such as random or combinatorial peptide or nonpeptide libraries which can be screened for molecules that specifically bind reproductive system related antigen polypeptides. Many libraries are known in the art

that can be used, e.g., chemically synthesized libraries, recombinant (e.g., phage display libraries), and *in vitro* translation-based libraries. Examples of chemically synthesized libraries are described in Fodor et al., 1991, Science 251:767-773; Houghten et al., 1991, Nature 354:84-86; Lam et al., 1991, Nature 354:82-84; Medynski, 1994, Bio/Technology 12:709-710;Gallop et al., 1994, J. Medicinal Chemistry 37(9):1233-1251; Ohlmeyer et al., 1993, Proc. Natl. Acad. Sci. USA 90:10922-10926; Erb et al., 1994, Proc. Natl. Acad. Sci. USA 91:11422-11426; Houghten et al., 1992, Biotechniques 13:412; Jayawickreme et al., 1994, Proc. Natl. Acad. Sci. USA 91:1614-1618; Salmon et al., 1993, Proc. Natl. Acad. Sci. USA 90:11708-11712; PCT Publication No. WO 93/20242; and Brenner and Lerner, 1992, Proc. Natl. Acad. Sci. USA 89:5381-5383.

- [0811] Examples of phage display libraries are described in Scott and Smith, 1990, Science 249:386-390; Devlin et al., 1990, Science, 249:404-406; Christian, R. B., et al., 1992, J. Mol. Biol. 227:711-718); Lenstra, 1992, J. Immunol. Meth. 152:149-157; Kay et al., 1993, Gene 128:59-65; and PCT Publication No. WO 94/18318 dated Aug. 18, 1994.
- [0812] In vitro translation-based libraries include but are not limited to those described in PCT Publication No. WO 91/05058 dated Apr. 18, 1991; and Mattheakis et al., 1994, Proc. Natl. Acad. Sci. USA 91:9022-9026.
- [0813] By way of examples of nonpeptide libraries, a benzodiazepine library (see e.g., Bunin et al., 1994, Proc. Natl. Acad. Sci. USA 91:4708-4712) can be adapted for use. Peptoid libraries (Simon et al., 1992, Proc. Natl. Acad. Sci. USA 89:9367-9371) can also be used. Another example of a library that can be used, in which the amide functionalities in peptides have been permethylated to generate a chemically transformed combinatorial library, is described by Ostresh et al. (1994, Proc. Natl. Acad. Sci. USA 91:11138-11142).
- [0814] The variety of non-peptide libraries that are useful in the present invention is great. For example, Ecker and Crooke, 1995, Bio/Technology 13:351-360 list benzodiazepines, hydantoins, piperazinediones, biphenyls, sugar analogs, beta-mercaptoketones, arylacetic acids, acylpiperidines, benzopyrans, cubanes, xanthines, aminimides, and oxazolones as among the chemical species that form the basis of various libraries.

[0815] Non-peptide libraries can be classified broadly into two types: decorated monomers and oligomers. Decorated monomer libraries employ a relatively simple scaffold structure upon which a variety functional groups is added. Often the scaffold will be a molecule with a known useful pharmacological activity. For example, the scaffold might be the benzodiazepine structure.

Non-peptide oligomer libraries utilize a large number of monomers that are assembled together in ways that create new shapes that depend on the order of the monomers. Among the monomer units that have been used are carbamates, pyrrolinones, and morpholinos. Peptoids, peptide-like oligomers in which the side chain is attached to the alpha amino group rather than the alpha carbon, form the basis of another version of non-peptide oligomer libraries. The first non-peptide oligomer libraries utilized a single type of monomer and thus contained a repeating backbone. Recent libraries have utilized more than one monomer, giving the libraries added flexibility.

Screening the libraries can be accomplished by any of a variety of commonly known methods. See, e.g., the following references, which disclose screening of peptide libraries: Parmley and Smith, 1989, Adv. Exp. Med. Biol. 251:215-218; Scott and Smith, 1990, Science 249:386-390; Fowlkes et al., 1992; BioTechniques 13:422-427; Oldenburg et al., 1992, Proc. Natl. Acad. Sci. USA 89:5393-5397; Yu et al., 1994, Cell 76:933-945; Staudt et al., 1988, Science 241:577-580; Bock et al., 1992, Nature 355:564-566; Tuerk et al., 1992, Proc. Natl. Acad. Sci. USA 89:6988-6992; Ellington et al., 1992, Nature 355:850-852; U.S. Pat. No. 5,096,815, U.S. Pat. No. 5,223,409, and U.S. Pat. No. 5,198,346, all to Ladner et al.; Rebar and Pabo, 1993, Science 263:671-673; and CT Publication No. WO 94/18318.

[0818] In a specific embodiment, screening to identify a molecule that binds reproductive system related antigen polypeptides can be carried out by contacting the library members with a reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides immobilized on a solid phase and harvesting those library members that bind to the reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides. Examples of such screening methods, termed "panning" techniques are described by way of example in Parmley and Smith, 1988, Gene 73:305-318; Fowlkes et al., 1992,

BioTechniques 13:422-427; International Publication No. WO 94/18318; and in references cited herein.

[0819] In another embodiment, the two-hybrid system for selecting interacting proteins in yeast (Fields and Song, 1989, Nature 340:245-246; Chien et al., 1991, Proc. Natl. Acad. Sci. USA 88:9578-9582) can be used to identify molecules that specifically bind to reproductive system related antigen polypeptides or reproductive system related antigen-like polypeptides.

[0820] Where the reproductive system related antigen binding molecule is a polypeptide, the polypeptide can be conveniently selected from any peptide library, including random peptide libraries, combinatorial peptide libraries, or biased peptide libraries. The term "biased" is used herein to mean that the method of generating the library is manipulated so as to restrict one or more parameters that govern the diversity of the resulting collection of molecules, in this case peptides.

[0821] Thus, a truly random peptide library would generate a collection of peptides in which the probability of finding a particular amino acid at a given position of the peptide is the same for all 20 amino acids. A bias can be introduced into the library, however, by specifying, for example, that a lysine occur every fifth amino acid or that positions 4, 8, and 9 of a decapeptide library be fixed to include only arginine. Clearly, many types of biases can be contemplated, and the present invention is not restricted to any particular bias. Furthermore, the present invention contemplates specific types of peptide libraries, such as phage displayed peptide libraries and those that utilize a DNA construct comprising a lambda phage vector with a DNA insert.

[0822] As mentioned above, in the case of a reproductive system related antigen binding molecule that is a polypeptide, the polypeptide may have about 6 to less than about 60 amino acid residues, preferably about 6 to about 10 amino acid residues, and most preferably, about 6 to about 22 amino acids. In another embodiment, a reproductive system related antigen binding polypeptide has in the range of 15-100 amino acids, or 20-50 amino acids.

[0823] The selected reproductive system related antigen binding polypeptide can be obtained by chemical synthesis or recombinant expression.

Other Activities

A polypeptide, polynucleotide, agonist, or antagonist of the present invention, as a result of the ability to stimulate vascular endothelial cell growth, may be employed in treatment for stimulating re-vascularization of ischemic tissues due to various disease conditions such as thrombosis, arteriosclerosis, and other cardiovascular conditions. The polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed to stimulate angiogenesis and limb regeneration, as discussed above.

[0825] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed for treating wounds due to injuries, burns, post-operative tissue repair, and ulcers since they are mitogenic to various cells of different origins, such as fibroblast cells and skeletal muscle cells, and therefore, facilitate the repair or replacement of damaged or diseased tissue.

[0826] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed to stimulate neuronal growth and to treat and prevent neuronal damage which occurs in certain neuronal disorders or neuro-degenerative conditions such as Alzheimer's disease, Parkinson's disease, and AIDS-related complex. A polypeptide, polynucleotide, agonist, or antagonist of the present invention may have the ability to stimulate chondrocyte growth, therefore, they may be employed to enhance bone and periodontal regeneration and aid in tissue transplants or bone grafts.

[0827] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may be also be employed to prevent skin aging due to sunburn by stimulating keratinocyte growth.

[0828] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed for preventing hair loss, since FGF family members activate hair-forming cells and promotes melanocyte growth. Along the same lines, a polypeptide, polynucleotide, agonist, or antagonist of the present invention may be employed to stimulate growth and differentiation of hematopoietic cells and bone marrow cells when used in combination with other cytokines.

[0829] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed to maintain organs before transplantation or for

supporting cell culture of primary tissues. A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed for inducing tissue of mesodermal origin to differentiate in early embryos.

[0830] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also increase or decrease the differentiation or proliferation of embryonic stem cells, besides, as discussed above, hematopoietic lineage.

[0831] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be used to modulate mammalian characteristics, such as body height, weight, hair color, eye color, skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cosmetic surgery). Similarly, a polypeptide, polynucleotide, agonist, or antagonist of the present invention may be used to modulate mammalian metabolism affecting catabolism, anabolism, processing, utilization, and storage of energy.

[0832] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may be used to change a mammal's mental state or physical state by influencing biorhythms, caricadic rhythms, depression (including depressive disorders), tendency for violence, tolerance for pain, reproductive capabilities (preferably by Activin or Inhibin-like activity), hormonal or endocrine levels, appetite, libido, memory, stress, or other cognitive qualities.

[0833] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be used as a food additive or preservative, such as to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors or other nutritional components.

[0834] The above-recited applications have uses in a wide variety of hosts. Such hosts include, but are not limited to, human, murine, rabbit, goat, guinea pig, camel, horse, mouse, rat, hamster, pig, micro-pig, chicken, goat, cow, sheep, dog, cat, non-human primate, and human. In specific embodiments, the host is a mouse, rabbit, goat, guinea pig, chicken, rat, hamster, pig, sheep, dog or cat. In preferred embodiments, the host is a mammal. In most preferred embodiments, the host is a human.

Other Preferred Embodiments

Other preferred embodiments of the claimed invention include an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.

- [0836] Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of the portion of SEQ ID NO:X as defined in column 4, "ORF (From-To)", in Table 1A.
- [0837] Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of the portion of SEQ ID NO:X as defined in columns 8 and 9, "NT From" and "NT To" respectively, in Table 2.
- [0838] Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 150 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.
- [0839] Further preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 500 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.
- [0840] A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of the portion of SEQ ID NO:X defined in column 4, "ORF (From-To)", in Table 1A.
- [0841] A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of the portion of SEQ ID NO:X defined in columns 8 and 9, "NT From" and "NT To", respectively, in Table 2.

[0842] A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.

- Also preferred is an isolated nucleic acid molecule which hybridizes under stringent hybridization conditions to a nucleic acid molecule comprising a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z, wherein said nucleic acid molecule which hybridizes does not hybridize under stringent hybridization conditions to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or of only T residues.
- [0844] Also preferred is a composition of matter comprising a DNA molecule which comprises the cDNA contained in Clone ID NO:Z.
- [0845] Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides of the cDNA sequence contained in Clone ID NO:Z.
- [0846] Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of an open reading frame sequence encoded by cDNA contained in Clone ID NO:Z.
- [0847] Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous nucleotides in the nucleotide sequence encoded by cDNA contained in Clone ID NO:Z.
- [0848] A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500 contiguous nucleotides in the nucleotide sequence encoded by cDNA contained in Clone ID NO:Z.
- [0849] A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by cDNA contained in Clone ID NO:Z.

sample a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence encoded by cDNA contained in Clone ID NO:Z; which method comprises a step of comparing a nucleotide sequence of at least one nucleic acid molecule in said sample with a sequence selected from said group and determining whether the sequence of said nucleic acid molecule in said selected sequence.

Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence determined from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

[0852] A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence of the cDNA contained in Clone ID NO:Z.

[0853] The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto; or the cDNA contained in Clone ID NO:Z which encodes a protein, wherein the method comprises a step of detecting in a biological sample obtained from said subject nucleic acid molecules, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence of cDNA contained in Clone ID NO:Z.

[0855] The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 4 of Table 1A or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence encoded by cDNA contained in Clone ID NO:Z. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

[0857] Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a DNA microarray or "chip" of at least 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 15, 20, 25, 30, 40, 50, 100, 150, 200, 250, 300, 500, 1000, 2000, 3000, or 4000 nucleotide sequences, wherein at least one sequence in said DNA microarray or "chip" is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected

from the group consisting of a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1A; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA "Clone ID" in Table 1A.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0859] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0860] Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0861] Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0862] Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a polypeptide encoded by contained in Clone ID NO:Z

[0863] Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a portion of said polypeptide encoded

by cDNA contained in Clone ID NO:Z; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or the polypeptide sequence of SEQ ID NO:Y.

[0864] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0865] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0866] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the amino acid sequence of a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0867] Further preferred is an isolated antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: a polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0868] Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: a polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z; which method comprises a step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group and determining whether the sequence of

said polypeptide molecule in said sample is at least 90% identical to said sequence of at least 10 contiguous amino acids.

Also preferred is the above method wherein said step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group comprises determining the extent of specific binding of polypeptides in said sample to an antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: a polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0870] Also preferred is the above method wherein said step of comparing sequences is performed by comparing the amino acid sequence determined from a polypeptide molecule in said sample with said sequence selected from said group.

Also preferred is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting polypeptide molecules in said sample, if any, comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0872] Also preferred is the above method for identifying the species, tissue or cell type of a biological sample, which method comprises a step of detecting polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the above group.

[0873] Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a nucleic acid sequence identified in Table 1A or Table 2 encoding a polypeptide, which method comprises a

step of detecting in a biological sample obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0874] In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding a polypeptide wherein said polypeptide comprises an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0876] Also preferred is an isolated nucleic acid molecule, wherein said nucleotide sequence encoding a polypeptide has been optimized for expression of said polypeptide in a prokaryotic host.

[0877] Also preferred is a polypeptide molecule, wherein said polypeptide comprises an amino acid sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0878] Further preferred is a method of making a recombinant vector comprising inserting any of the above isolated nucleic acid molecule into a vector. Also preferred is the recombinant vector produced by this method. Also preferred is a method of making a recombinant host cell comprising introducing the vector into a host cell, as well as the recombinant host cell produced by this method.

Also preferred is a method of making an isolated polypeptide comprising culturing this recombinant host cell under conditions such that said polypeptide is expressed and recovering said polypeptide. Also preferred is this method of making an isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said polypeptide is a human protein comprising an amino acid sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z. The isolated polypeptide produced by this method is also preferred.

[0880] Also preferred is a method of treatment of an individual in need of an increased level of a protein activity, which method comprises administering to such an individual a Therapeutic comprising an amount of an isolated polypeptide, polynucleotide, immunogenic fragment or analogue thereof, binding agent, antibody, or antigen binding fragment of the claimed invention effective to increase the level of said protein activity in said individual.

[0881] Also preferred is a method of treatment of an individual in need of a decreased level of a protein activity, which method comprised administering to such an individual a Therapeutic comprising an amount of an isolated polypeptide, polynucleotide, immunogenic fragment or analogue thereof, binding agent, antibody, or antigen binding fragment of the claimed invention effective to decrease the level of said protein activity in said individual.

[0882] Also preferred is a method of treatment of an individual in need of a specific delivery of toxic compositions to diseased cells (e.g., tumors, leukemias or lymphomas), which method comprises administering to such an individual a Therapeutic comprising an amount of an isolated polypeptide of the invention, including, but not limited to a binding agent, or antibody of the claimed invention that are associated with toxin or cytotoxic prodrugs.

[0883] Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

TABLE 6

ATCC Deposits	Deposit Date	ATCC Designation Number
LP01, LP02, LP03,	May-20-97	209059, 209060, 209061, 209062, 209063,
LP04, LP05, LP06,		209064, 209065, 209066, 209067, 209068,
LP07, LP08, LP09,	1	209069
LP10, LP11,		
LP12	Jan-12-98	209579
LP13	Jan-12-98	209578
LP14	Jul-16-98	203067
LP15	Jul-16-98	203068
LP16	Feb-1-99	203609
LP17	Feb-1-99	203610
LP20	Nov-17-98	203485
LP21	Jun-18-99	PTA-252
LP22	Jun-18-99	PTA-253
LP23	Dec-22-99	PTA-1081

Examples

Example 1: Isolation of a Selected cDNA Clone From the Deposited Sample

[0884] Each Clone ID NO:Z is contained in a plasmid. Table 7 identifies the vectors used to construct the cDNA library from which each clone was isolated. In many cases, the vector used to construct the library is a phage vector from which a plasmid has been excised. The following correlates the related plasmid for each phage vector used in constructing the cDNA library. For example, where a particular clone is identified in Table 7 as being isolated in the vector "Lambda Zap," the corresponding deposited clone is in "pBluescript."

Vector Used to Construct Library	Corresponding Deposited Plasmid
Lambda Zap	pBluescript (pBS)
Uni-Zap XR	pBluescript (pBS)
Zap Express	pBK
lafmid BA	plafmid BA
pSport1	pSport1
pCMVSport 2.0	pCMVSport 2.0
pCMVSport 3.0	pCMVSport 3.0
pCR®2.1	pCR [®] 2.1

Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., Nucleic Acids Res. 16:7583-7600 (1988); Alting-Mees, M. A. and Short, J. M., Nucleic Acids Res. 17:9494 (1989)) and pBK (Alting-Mees, M. A. et al., Strategies 5:58-61 (1992)) are commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK contains a neomycin resistance gene. Both can be transformed into E. coli strain XL-1 Blue, also available from Stratagene. pBS comes in 4 forms SK+, SK-, KS+ and KS. The S and K refers to the orientation of the polylinker to the T7 and T3 primer

sequences which flank the polylinker region ("S" is for SacI and "K" is for Kpnl which are the first sites on each respective end of the linker). "+" or "-" refer to the orientation of the fl origin of replication ("ori"), such that in one orientation, single stranded rescue initiated from the fl ori generates sense strand DNA and in the other, antisense.

Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into E. coli strain DH10B, also available from Life Technologies. (See, for instance, Gruber, C. E., et al., Focus 15:59 (1993).) Vector lafmid BA (Bento Soares, Columbia University, NY) contains an ampicillin resistance gene and can be transformed into E. coli strain XL-1 Blue. Vector pCR®2.1, which is available from Invitrogen, 1600 Faraday Avenue, Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed into E. coli strain DH10B, available from Life Technologies. (See, for instance, Clark, J. M., Nuc. Acids Res. 16:9677-9686 (1988) and Mead, D. et al., Bio/Technology 9: (1991).) Preferably, a polynucleotide of the present invention does not comprise the vector sequences identified for the particular clone in Table 7, as well as the corresponding plasmid vector sequences designated above.

[0887] The deposited material in the sample assigned the ATCC Deposit Number cited by reference to Tables 1A, 2, 6 and 7 for any given cDNA clone also may contain one or more additional plasmids, each comprising a cDNA clone different from that given clone. Thus, deposits sharing the same ATCC Deposit Number contain at least a plasmid for each Clone ID NO:Z.

TABLE 7

Libraries owned by Catalog	Catalog Description	Vector	ATCC
			Deposit
HUKA HUKB HUKC HUKD	Human Uterine Cancer	Lambda ZAP II	LP01
HUKE HUKF HUKG	1		
HCNA HCNB	Human Colon	Lambda Zap II	LP01
HFFA	Human Fetal Brain, random primed	Lambda Zap II	LP01
HTWA	Resting T-Cell	Lambda ZAP II	LP01

Libraries owned by Catalog	Catalog Description	Vector	ATCC ·
		,	Deposit
НВQА	Early Stage Human Brain, random primed	Lambda ZAP II	LP01
HLMB HLMF HLMG HLMH	breast lymph node CDNA library	Lambda ZAP II	LP01
НГМІ НГМІ НГММ НГМИ			
HCQA HCQB	human colon cancer	Lamda ZAP II	LP01
HMEA HMEC HMED HMEE	Human Microvascular Endothelial	Lambda ZAP II	LP01
нмег нмес нмеі нмеј	Cells, fract. A		,
HMEK HMEL			
HUSA HUSC	Human Umbilical Vein Endothelial	Lambda ZAP II	LP01
·	Cells, fract. A	1	
HLQA HLQB	Hepatocellular Tumor	Lambda ZAP II	LP01
HHGA HHGB HHGC HHGD	Hemangiopericytoma	Lambda ZAP II	LP01
HSDM	Human Striatum Depression, re-rescue	Lambda ZAP II	LP01
HUSH	H Umbilical Vein Endothelial Cells,	Lambda ZAP II	LP01
	frac A, re-excision		
HSGS	Salivary gland, subtracted	Lambda ZAP II	LP01
HFXA HFXB HFXC HFXD	Brain frontal cortex	Lambda ZAP II	LP01
HFXE HFXF HFXG HFXH	·		
НРОА НРОВ НРОС	PERM TF274	Lambda ZAP II	LP01
HFXJ HFXK	Brain Frontal Cortex, re-excision	Lambda ZAP II	LP01
HCWA HCWB HCWC HCWD	CD34 positive cells (Cord Blood)	ZAP Express	LP02
HCWE HCWF HCWG HCWH			
HCWI HCWJ HCWK			
HCUA HCUB HCUC	CD34 depleted Buffy Coat (Cord	ZAP Express	LP02
	Blood)		
HRSM	A-14 cell line	ZAP Express	LP02
HRSA	A1-CELL LINE	ZAP Express	LP02
HCUD HCUE HCUF HCUG	CD34 depleted Buffy Coat (Cord	ZAP Express	LP02
нсин нсиј	Blood), re-excision		
HBXE HBXF HBXG	H. Whole Brain #2, re-excision	ZAP Express	LP02
HRLM	L8 cell line	ZAP Express	LP02
HBXA HBXB HBXC HBXD	Human Whole Brain #2 - Oligo dT >	ZAP Express	LP02
HUDA HUDB HUDC	Testes ·	ZAP Express	LP02
НТМ ННТО	H. hypothalamus, frac A;re-excision	ZAP Express	LP02
HTL	H. hypothalamus, frac A	ZAP Express	LP02
IASA HASD	Human Adult Spleen	Uni-ZAP XR	LP03
IFKC HFKD HFKE HFKF	Human Fetal Kidney	Uni-ZAP XR	LP03
IFKG	· [

Libraries owned by Catalog	Catalog Description	Vector	ATCC
		.	Deposit
HE8A HE8B HE8C HE8D HE8E	Human 8 Week Whole Embryo	Uni-ZAP XR	LP03
HE8F HE8M HE8N			
HGBA HGBD HGBE HGBF	Human Gall Bladder	Uni-ZAP XR	LP03
ндво новн нові			
HLHA HLHB HLHC HLHD	Human Fetal Lung III	Uni-ZAP XR	LP03
нгне нгнь нгно нгнн			
нгнб			
HPMA HPMB HPMC HPMD	Human Placenta	Uni-ZAP XR	LP03
НРМЕ НРМГ НРМС НРМН			
HPRA HPRB HPRC HPRD	Human Prostate	Uni-ZAP XR	LP03
HSIA HSIC HSID HSIE	Human Adult Small Intestine	Uni-ZAP XR	LP03
HTEA HTEB HTEC HTED HTEE		Uni-ZAP XR	LP03
НТЕГ НТЕС НТЕН НТЕІ НТЕЈ			2.05
НТЕК			
НТРА НТРВ НТРС НТРО НТРЕ	Human Pancreas Tumor	Uni-ZAP XR	LP03
HTTA HTTB HTTC HTTD HTTE		Uni-ZAP XR	LP03
HTTF		OM-ZAF AK	LPUS
НАРА НАРВ НАРС НАРМ	Human Adult Pulmonary	Uni-ZAP XR	LP03
НЕТА НЕТВ НЕТС НЕТО НЕТЕ	Human Endometrial Tumor	Uni-ZAP XR	LP03
нет т нето нетн неті	·		
HHFB HHFC HHFD HHFE HHFF	Human Fetal Heart	Uni-ZAP XR	LP03
ннгс ннгн ннгі			
ННРВ ННРС ННРО ННРЕ ННРF	Human Hippocampus	Uni-ZAP XR	LP03
ННРС ННРН			
HCE1 HCE2 HCE3 HCE4 HCE5	Human Cerebellum	Uni-ZAP XR	LP03
HCEB HCEC HCED HCEE HCEF			1
HCEG	, ,		
HUVB HUVC HUVD HUVE	Human Umbilical Vein, Endo. remake	Uni-ZAP XR	LP03
HSTA HSTB HSTC HSTD	Human Skin Tumor	Uni-ZAP XR	LP03
TAA HTAB HTAC HTAD	Human Activated T-Cells	Uni-ZAP XR	LP03
ITAE			2. 03
IFEA HFEB HFEC	Human Fetal Epithelium (Skin)	Uni-ZAP XR	LP03
НРА НЈРВ НЈРС НЈРD	HUMAN JURKAT MEMBRANE	Uni-ZAP XR	LP03
i	BOUND POLYSOMES		
IESA	Human epithelioid sarcoma	Uni-Zap XR	LP03
ILTA HLTB HLTC HLTD HLTE	Human T-Cell Lymphoma	Uni-ZAP XR	LP03
ILTF			
IFTA HFTB HFTC HFTD	Human Fetal Dura Mater	Uni-ZAP XR	LP03
	Human Rhabdomyosarcoma	Uni-ZAP XR	LP03

Libraries owned by Catalog	Catalog Description	Vector	IATCC
		,	Deposit
HRDE HRDF			
НСАА НСАВ НСАС	Cem cells cyclohexamide treated	Uni-ZAP XR	LP03
HRGA HRGB HRGC HRGD	Raji Cells, cyclohexamide treated	Uni-ZAP XR	LP03
HSUA HSUB HSUC HSUM	Supt Cells, cyclohexamide treated	Uni-ZAP XR	LP03
HT4A HT4C HT4D	Activated T-Cells, 12 hrs.	Uni-ZAP XR	LP03
НЕ9А НЕ9В НЕ9С НЕ9D НЕ9Е	Nine Week Old Early Stage Human	Uni-ZAP XR	LP03
не9f не9g не9h не9м не9h			2.03
HATA HATB HATC HATD	Human Adrenal Gland Tumor	Uni-ZAP XR	LP03
НАТЕ			
НТ5А	Activated T-Cells, 24 hrs.	Uni-ZAP XR	LP03
HFGA HFGM	Human Fetal Brain	Uni-ZAP XR	LP03
HNEA HNEB HNEC HNED	Human Neutrophil	Uni-ZAP XR	LP03
HNEE			2.03
HBGB HBGD	Human Primary Breast Cancer	Uni-ZAP XR	LP03
HBNA HBNB	Human Normal Breast	Uni-ZAP XR	LP03
HCAS	Cem Cells, cyclohexamide treated,	Uni-ZAP XR	LP03
	subtra		El 05
HHPS	Human Hippocampus, subtracted	pBS	LP03
HKCS HKCU	Human Colon Cancer, subtracted	pBS	LP03
HRGS	Raji cells, cyclohexamide treated,	pBS	LP03
	subtracted	ľ	
HSUT	Supt cells, cyclohexamide treated,	pBS	LP03
	differentially expressed	ľ	
HT4S	Activated T-Cells, 12 hrs, subtracted	Uni-ZAP XR	LP03
HCDA HCDB HCDC HCDD	Human Chondrosarcoma	Uni-ZAP XR	LP03
HCDE			
HOAA HOAB HOAC	Human Osteosarcoma	Uni-ZAP XR	LP03
HTLA HTLB HTLC HTLD HTLE	Human adult testis, large inserts	Uni-ZAP XR	LP03
HTLF		Į	
HLMA HLMC HLMD	Breast Lymph node cDNA library	Uni-ZAP XR	LP03
Н6ЕА Н6ЕВ Н6ЕС	HL-60, PMA 4H	Uni-ZAP XR	LP03
ITXA HTXB HTXC HTXD	Activated T-Cell (12hs)/Thiouridine	Uni-ZAP XR	LP03
ITXE HTXF HTXG HTXH	labelledEco		
INFA HNFB HNFC HNFD	Human Neutrophil, Activated	Uni-ZAP XR	LP03
INFE HNFF HNFG HNFH HNFJ			
тов нтос	HUMAN TONSILS, FRACTION 2	Uni-ZAP XR	LP03
IMGB	Human OB MG63 control fraction I	Uni-ZAP XR	LP03
IOPB	Human OB HOS control fraction I	Uni-ZAP XR	LP03
IORB	Human OB HOS treated (10 nM E2)	Uni-ZAP XR	LP03

Libraries owned by Catalog	Catalog Description	Vector	ATCC
, ,		Vector	ATCC
	fraction I		Deposit
HSVA HSVB HSVC	Human Chronic Synovitis	Uni-ZAP XR	1, 202
HROA	HUMAN STOMACH	Uni-ZAP XR	LP03
НВЈА НВЈВ НВЈС НВЈО НВЈЕ	HUMAN B CELL LYMPHOMA	Uni-ZAP XR	LP03
НВЈҒ НВЈС НВЈН НВЈІ НВЈЈ	TOWN B CEEL ET WIT HOWA	Om-ZAP XR	LP03
НВЈК	,		
HCRA HCRB HCRC	human corpus colosum	Uni-ZAP XR	1 000
HODA HODB HODC HODD	human ovarian cancer	Uni-ZAP XR	LP03
HDSA	Dermatofibrosarcoma Protuberance	Uni-ZAP XR	LP03
HMWA HMWB HMWC HMWD	Bone Marrow Cell Line (RS4;11)	Uni-ZAP XR	LP03
HMWE HMWF HMWG HMWH	Zone Mariow Cell Line (R34,11)	Uni-ZAP XR	LP03
HMWI HMWJ	,		
HSOA	stomach cancer (human)	Uni-ZAP XR	7 702
HERA	SKIN	Uni-ZAP XR	LP03
HMDA	Brain-medulloblastoma	Uni-ZAP XR	LP03
HGLA HGLB HGLD	Glioblastoma	Uni-ZAP XR	LP03
HEAA	H. Atrophic Endometrium	Uni-ZAP XR	LP03
НВСА НВСВ	H. Lymph node breast Cancer		LP03
IPWT	Human Prostate BPH, re-excision	Uni-ZAP XR	LP03
IFVG HFVH HFVI	Fetal Liver, subtraction II	Uni-ZAP XR	LP03
INFI	Human Neutrophils, Activated, re-	pBS	LP03
	lexcision	pBS	LP03
IBMB HBMC HBMD	Human Bone Marrow, re-excision	pBS	LP03
IKML HKMM HKMN	H. Kidney Medulla, re-excision	pBS	LP03
IKIX HKIY	H. Kidney Cortex, subtracted	pBS	LP03
IADT	H. Amygdala Depression, subtracted	pBS	LP03
6AS	HI-60, untreated, subtracted	Uni-ZAP XR	LP03
6ES	HL-60, PMA 4H, subtracted	Uni-ZAP XR	1
CDC .	HL-60, RA 4h, Subtracted		LP03
	HL-60, PMA 1d, subtracted	Uni-ZAP XR	LP03
	Activated T-cell(12h)/Thiouridine-re-	Uni-ZAP XR	LP03
	excision	Uni-ZAP XR	LP03
	Monocyte activated	II-: ZADAD	
MSE HMSF HMSG HMSH	Monocyte activated	Uni-ZAP XR	LP03
MSI HMSJ HMSK			
	Human Amygdala	lini ZAR VR	V 700
AGE HAGF		Uni-ZAP XR	LP03
	STROMAL -OSTEOCLASTOMA	Ilm: 7AD NO	-
	- WOMEN TO LECTORAS LOIMIN	Uni-ZAP XR	LP03

Libraries owned by Catalog	Catalog Description	Vector	ATCC
			Deposit
	unamplified		
HSQA HSQB HSQC HSQD	Stromal cell TF274	Uni-ZAP XR	LP03
HSQE HSQF HSQG			
HSKA HSKB HSKC HSKD	Smooth muscle, serum treated	Uni-ZAP XR	LP03
HSKE HSKF HSKZ			
HSLA HSLB HSLC HSLD HSLE	Smooth muscle,control	Uni-ZAP XR	LP03
HSLF HSLG			
HSDA HSDD HSDE HSDF	Spinal cord	Uni-ZAP XR	LP03
HSDG HSDH	·		
HPWS	Prostate-BPH subtracted II	pBS	LP03
HSKW HSKX HSKY	Smooth Muscle-HASTE normalized	pBS	LP03
HFPB HFPC HFPD	H. Frontal cortex,epileptic;re-excision	Uni-ZAP XR	LP03
HSDI HSDJ HSDK	Spinal Cord, re-excision	Uni-ZAP XR	LP03
HSKN HSKO	Smooth Muscle Serum Treated, Norm	pBS	LP03
HSKG HSKH HSKI	Smooth muscle, serum induced,re-exc	pBS	LP03
HFCA HFCB HFCC HFCD HFC	· ·	Uni-ZAP XR	
HFCF	Januari Tetar Brani	UIII-ZAP AR	LP04
HPTA HPTB HPTD	Human Pituitary	Uni-ZAP XR	LP04
НТНВ НТНС HTHD	Human Thymus	Uni-ZAP XR	LP04
HE6B HE6C HE6D HE6E HE6F	Human Whole Six Week Old Embryo	Uni-ZAP XR	LP04
HE6G HE6S	·		
HSSA HSSB HSSC HSSD HSSE	Human Synovial Sarcoma	Uni-ZAP XR	LP04
HSSF HSSG HSSH HSSI HSSJ			
HSSK			
HE7T	7 Week Old Early Stage Human,	Uni-ZAP XR	LP04
	subtracted		
НЕРА НЕРВ НЕРС	Human Epididymus	Uni-ZAP XR	LP04
HSNA HSNB HSNC HSNM	Human Synovium	Uni-ZAP XR	LP04
HSNN			
HPFB HPFC HPFD HPFE	Human Prostate Cancer, Stage C	Uni-ZAP XR	LP04
	fraction		
HE2A HE2D HE2E HE2H HE2I	12 Week Old Early Stage Human	Uni-ZAP XR	LP04
HE2M HE2N HE2O			
HE2B HE2C HE2F HE2G HE2P	12 Week Old Early Stage Human, II	Uni-ZAP XR	LP04
łE2Q		1	
ІРТЅ НРТТ НРТИ	Human Pituitary, subtracted	Uni-ZAP XR	LP04
IAUA HAUB HAUC	Amniotic Cells - TNF induced	Uni-ZAP XR	LP04
IAQA HAQB HAQC HAQD	Amniotic Cells - Primary Culture	Uni-ZAP XR	LP04
IWTA HWTB HWTC	wilm's tumor	Uni-ZAP XR	LP04

Libraries owned by Catalog	Catalog Description	Vector	ATCC
Lioranes owned by Catalog	Catalog Description	Vector	
HBSD	Des Constitution	11:715.45	Deposit
HSGB	Bone Cancer, re-excision	Uni-ZAP XR	LP04
	Salivary gland, re-excision	Uni-ZAP XR	LP04
HSJA HSJB HSJC	Smooth muscle-ILb induced	Uni-ZAP XR	LP04
HSXA HSXB HSXC HSXD	Human Substantia Nigra	Uni-ZAP XR	LP04
HSHA HSHB HSHC	Smooth muscle, IL1b induced	Uni-ZAP XR	LP04
HOUA HOUB HOUC HOUD	Adipocytes	Uni-ZAP XR	LP04
HOUE ,			
HPWA HPWB HPWC HPWD	Prostate BPH	Uni-ZAP XR	LP04
HPWE .			
HELA HELB HELC HELD HELI	E Endothelial cells-control	Uni-ZAP XR	LP04
HELF HELG HELH			
HEMA HEMB HEMC HEMD	Endothelial-induced	Uni-ZAP XR	LP04
НЕМЕ НЕМГ НЕМС НЕМН			
HBIA HBIB HBIC	Human Brain, Striatum	Uni-ZAP XR	LP04
HHSA HHSB HHSC HHSD	Human Hypothalmus, Schizophrenia	Uni-ZAP XR	LP04
HHSE			
HNGA HNGB HNGC HNGD	neutrophils control	Uni-ZAP XR	LP04
HNGE HNGF HNGG HNGH	1 .		
HNGI HNGJ		}	
HNHA HNHB HNHC HNHD	Neutrophils IL-1 and LPS induced	Uni-ZAP XR	LP04
HNHE HNHF HNHG HNHH			
НИНІ НИНЈ			
HSDB HSDC	STRIATUM DEPRESSION	Uni-ZAP XR	LP04
ННРТ	Hypothalamus	Uni-ZAP XR	LP04
HSAT HSAU HSAV HSAW	Anergic T-cell	Uni-ZAP XR	LP04
HSAX HSAY HSAZ	, and good a con-		21 04
HBMS HBMT HBMU HBMV	Bone marrow	Uni-ZAP XR	LP04
НВМW НВМХ			2.04
HOEA HOEB HOEC HOED	Osteoblasts	Uni-ZAP XR	LP04
HOEE HOEF HOEJ	00.000.000	OIII-ZAII AR	L. 04
HAIA HAIB HAIC HAID HAIE	Epithelial-TNFa and INF induced	Uni-ZAP XR	LP04
HAIF	Dymichai- 1111 a and 1111 madeed	OIII-ZAI XX	LF 04
HTGA HTGB HTGC HTGD	Apoptotic T-cell	Uni-ZAP XR	LP04
HMCA HMCB HMCC HMCD	1		
HMCE	Macrophage-oxLDL	Uni-ZAP XR	LP04
	Managhay (CN) COT : 12	The GARAGE	li no :
HMAA HMAB HMAC HMAD	Macrophage (GM-CSF treated)	Uni-ZAP XR	LP04
HMAE HMAF HMAG	N. 10		
НРНА	Normal Prostate	Uni-ZAP XR	LP04
HPIA HPIB HPIC	LNCAP prostate cell line	Uni-ZAP XR	LP04

Libraries owned by Catalog	Catalog Description	Vector	ATCC
			Deposit
НРЈА НРЈВ НРЈС	PC3 Prostate cell line	Uni-ZAP XR	LP04
HOSE HOSF HOSG	Human Osteoclastoma, re-excision	Uni-ZAP XR	LP04
HTGE HTGF	Apoptotic T-cell, re-excision	Uni-ZAP XR	LP04
HMAJ HMAK	H Macrophage (GM-CSF treated), re- excision	Uni-ZAP XR	LP04
HACB HACC HACD	Human Adipose Tissue, re-excision	Uni-ZAP XR	LP04
НГРА	H. Frontal Cortex, Epileptic	Uni-ZAP XR	LP04
HFAA HFAB HFAC HFAD HFAE	Alzheimers, spongy change	Uni-ZAP XR	LP04
HFAM	Frontal Lobe, Dementia	Uni-ZAP XR	LP04
НМІА НМІВ НМІС	Human Manic Depression Tissue	Uni-ZAP XR	LP04
HTSA HTSE HTSF HTSG HTSH	Human Thymus	pBS	LP05
НРВА НРВВ НРВС НРВО НРВЕ	Human Pineal Gland	pBS	LP05
HSAA HSAB HSAC	HSA 172 Cells	pBS	LP05
HSBA HSBB HSBC HSBM	HSC172 cells	pBS	LP05
HJAA HJAB HJAC HJAD	Jurkat T-cell G1 phase	pBS	LP05
НЈВА НЈВВ НЈВС НЈВD	Jurkat T-Cell, S phase	pBS	LP05
HAFA HAFB	Aorta endothelial cells + TNF-a	pBS	LP05
HAWA HAWB HAWC	Human White Adipose	pBS	LP05
HTNA HTNB	Human Thyroid	pBS	LP05
ANOH	Normal Ovary, Premenopausal	pBS	LP05
HARA HARB	Human Adult Retina	pBS	LP05
ILJA HLJB	Human Lung	pCMVSport 1	LP06
НОРМ НОРО	H. Ovarian Tumor, II, OV5232	pCMVSport 2.0	LP07
HOGA HOGB HOGC	OV 10-3-95	pCMVSport 2.0	LP07
HCGL	CD34+cells, II	pCMVSport 2.0	LP07
IDLA	Hodgkin's Lymphoma I	pCMVSport 2.0	LP07
IDTA HDTB HDTC HDTD IDTE	Hodgkin's Lymphoma II	pCMVSport 2.0	LP07
KAA HKAB HKAC HKAD	Keratinocyte	pCMVSport2.0	LP07
IKAE HKAF HKAG HKAH		-	
ICIM	CAPFINDER, Crohn's Disease, lib 2	pCMVSport 2.0	LP07
IKAL	Keratinocyte, lib 2	pCMVSport2.0	LP07
IKAT	Keratinocyte, lib 3	pCMVSport2.0	LP07
INDA	Nasal polyps	pCMVSport2.0	LP07
IDRA	H. Primary Dendritic Cells,lib 3	pCMVSport2.0	LP07
ОНА НОНВ НОНС	Human Osteoblasts II	pCMVSport2.0	LP07
LDA HLDB HLDC	Liver, Hepatoma	pCMVSport3.0	LP08

Libraries owned by Catalog	Catalog Description	Vector	ATCC
		,	Deposit
HLDN HLDO HLDP	Human Liver, normal	pCMVSport3.0	LP08
НМТА	pBMC stimulated w/ poly I/C	pCMVSport3.0	LP08
HNTA	NTERA2, control	pCMVSport3.0	LP08
HDPA HDPB HDPC HDPD	Primary Dendritic Cells, lib !	pCMVSport3.0	LP08
HDPF HDPG HDPH HDPI HDF	P.I		
HDPK	l'		
HDPM HDPN HDPO HDPP	Primary Dendritic cells,frac 2	pCMVSport3.0	LP08
НМИА НМИВ НМИС	Myoloid Progenitor Cell Line	pCMVSport3.0	LP08
ННЕА ННЕВ ННЕС HHED	T Cell helper I	pCMVSport3.0	LP08
ННЕМ ННЕМ ННЕО ННЕР	T cell helper II	pCMVSport3.0	LP08
HEQA HEQB HEQC	Human endometrial stromal cells	pCMVSport3.0	LP08
НЈМА НЈМВ	Human endometrial stromal cells-	pCMVSport3.0	LP08
	treated with progesterone	, ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	12. 00
HSWA HSWB HSWC	Human endometrial stromal cells-	pCMVSport3.0	LP08
	treated with estradiol	,	2.00
HSYA HSYB HSYC	Human Thymus Stromal Cells	pCMVSport3.0	LP08
HLWA HLWB HLWC	Human Placenta	pCMVSport3.0	LP08
HRAA HRAB HRAC	Rejected Kidney, lib 4	pCMVSport3.0	LP08
НМТМ	PCR, pBMC I/C treated	PCRII	LP09
НМЈА	H. Meniingima, M6	pSport 1	LP10
НМКА НМКВ НМКС НМКD	H. Meningima, MI	pSport 1	LP10
НМКЕ	1	,	12.10
HUSG HUSI	Human umbilical vein endothelial cells,	pSport 1	LP10
	IL-4 induced		
HUSX HUSY	Human Umbilical Vein Endothelial	pSport 1	LP10
	Cells, uninduced		12.10
IOFA	Ovarian Tumor I, OV5232	pSport I	LP10
ICFA HCFB HCFC HCFD	T-Cell PHA 16 hrs	pSport 1	LP10
ICFL HCFM HCFN HCFO	T-Cell PHA 24 hrs	pSport I	LP10
ADA HADC HADD HADE	Human Adipose	pSport 1	LP10
ADF HADG			Di io
OVA HOVB HOVC	Human Ovary	pSport I	LP10
TWB HTWC HTWD HTWE	Resting T-Cell Library,II	pSport 1	LP10
TWF	3,4	popoli i	LF10
MMA	Spleen metastic melanoma	pSport 1	LP10
LYA HLYB HLYC HLYD	Spleen, Chronic lymphocytic leukemia	pSport I	
LYE	, suproof to tourchild	Popoli I	LP10
CGA	CD34+ cell, J	pSport 1	LP10
	, · · · · · · · · · · · · · · · · · · ·	Popore 1	ILTIU

Carlos Description	122	
	Vector	ATCC .
	<u> </u>	Deposit
		LP10
		LP10
	<u>' ' </u>	LP10
	pSport 1	LP10
Crohn's Disease	pSport I	LP10
HEL cell line	pSport 1	LP10
Human Astrocyte	pSport 1	LP10
Ulcerative Colitis	pSport I	LP10
NTERA2 + retinoic acid, 14 days	pSport I	LP10
Primary Dendritic cells, CapFinder2,	pSport 1	LP10
frac 1		
Primary Dendritic Cells, CapFinder,	pSport 1	LP10
frac 2		
Human Liver, normal,	pSport 1	LPIO
Human Dermal Endothelial	pSport1	LP10
Cells,untreated		
Human Dermal Endothelial cells,treated	pSport1	LP10
Human Stromal Endometrial	pSport1	LP10
fibroblasts, untreated	!	
Human Stromal endometrial fibroblasts,	pSport1	LP10
treated w/ estradiol		:
Human Stromal endometrial fibroblasts,	pSport1	LP10
treated with progesterone		
Human ovary tumor cell OV350721	pSport1	LP10
- 	1	
Merkel Cells	pSport1	LP10
Merkel Cells Pancreas Islet Cell Tumor	pSport1 pSport1	LP10 LP10
	· ·	
Pancreas Islet Cell Tumor	pSport1	LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2	pSport1 pSport1	LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2	pSport1 pSport1 pSport 1	LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate, BPH, Lib 2 Prostate BPH, Lib 2, subtracted	pSport1 pSport1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2 Prostate BPH,Lib 2, subtracted Synovial Fibroblasts (control)	pSport1 pSport1 pSport 1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2 Prostate BPH,Lib 2, subtracted Synovial Fibroblasts (control) Synovial hypoxia Synovial IL-1/TNF stimulated	pSport1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10 LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2 Prostate BPH,Lib 2, subtracted Synovial Fibroblasts (control) Synovial hypoxia Synovial IL-1/TNF stimulated Messangial cell, frac 1	pSport1 pSport1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10 LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2 Prostate BPH,Lib 2, subtracted Synovial Fibroblasts (control) Synovial hypoxia Synovial IL-1/TNF stimulated Messangial cell, frac 1 Bone Marrow Stromal Cell, untreated	pSport1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10 LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2 Prostate BPH,Lib 2, subtracted Synovial Fibroblasts (control) Synovial hypoxia Synovial IL-1/TNF stimulated Messangial cell, frac 1 Bone Marrow Stromal Cell, untreated Synovial Fibroblasts (II1/TNF), subt	pSport1 pSport1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10 LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2 Prostate BPH,Lib 2, subtracted Synovial Fibroblasts (control) Synovial hypoxia Synovial IL-1/TNF stimulated Messangial cell, frac 1 Bone Marrow Stromal Cell, untreated Synovial Fibroblasts (III/TNF), subt Synovial hypoxia-RSF subtracted	pSport1 pSport1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10 LP10 LP10 LP10
Pancreas Islet Cell Tumor Skin, burned Prostate,BPH, Lib 2 Prostate BPH,Lib 2, subtracted Synovial Fibroblasts (control) Synovial hypoxia Synovial IL-1/TNF stimulated Messangial cell, frac 1 Bone Marrow Stromal Cell, untreated Synovial Fibroblasts (II1/TNF), subt Synovial hypoxia-RSF subtracted Human Activated Monocytes	pSport1 pSport1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1 pSport 1	LP10 LP10 LP10 LP10 LP10 LP10 LP10 LP10
	Human Astrocyte Ulcerative Colitis NTERA2 + retinoic acid, 14 days Primary Dendritic cells, CapFinder2, frac 1 Primary Dendritic Cells, CapFinder, frac 2 Human Liver, normal, Human Dermal Endothelial Cells, untreated Human Stromal Endometrial fibroblasts, untreated Human Stromal endometrial fibroblasts, treated w/ estradiol Human Stromal endometrial fibroblasts, treated with progesterone	Human Tonsil, Lib 3 pSport 1 Salivary Gland, Lib 2 pSport 1 Breast Cancer cell line, MDA 36 pSport 1 Breast Cancer Cell line, angiogenic pSport 1 Crohn's Disease pSport 1 HEL cell line pSport 1 Human Astrocyte pSport 1 Ulcerative Colitis pSport 1 NTERA2 + retinoic acid, 14 days pSport 1 Primary Dendritic cells, CapFinder 2, pSport 1 Primary Dendritic Cells, CapFinder, pSport 1 Primary Dendritic Cells, CapFinder, pSport 1 Human Liver, normal, pSport 1 Human Dermal Endothelial pSport 1 Cells, untreated Human Stromal Endometrial fibroblasts, untreated Human Stromal endometrial fibroblasts, pSport 1 treated w/ estradiol Human Stromal endometrial fibroblasts, pSport 1 treated with progesterone

Libraries owned by Catalog	Catalog Description	Vector	ATCC
			Deposit
ННВЕ			
НВВА НВВВ	Human Brain	pCMVSport 1	LP012
HLJA HLJB HLJC HLJD HLJE	Human Lung	pCMVSport 1	LP012
HOGA HOGB HOGC	Ovarian Tumor	pCMVSport 2.0	LP012
НТЈМ	Human Tonsils, Lib 2	pCMVSport 2.0	LP012
HAMF HAMG	КМН2	pCMVSport 3.0	LP012
НАЈА НАЈВ НАЈС	L428	pCMVSport 3.0	LP012
HWBA HWBB HWBC HWBD HWBE	Dendritic cells, pooled	pCMVSport 3.0	LP012
HWAA HWAB HWAC HWAD HWAE	Human Bone Marrow, treated	pCMVSport 3.0	LP012
НУАА НУАВ НУАС	B Cell lymphoma	pCMVSport 3.0	LP012
нwнg нwнн нwні	Healing groin wound, 6.5 hours post incision	pCMVSport 3.0	LP012
нwнр нwнQ нwнr	Healing groin wound; 7.5 hours post incision	pCMVSport 3.0	LP012
HARM	Healing groin wound - zero hr post- incision (control)	pCMVSport 3.0	LP012
НВІМ	Olfactory epithelium; nasalcavity	pCMVSport 3.0	LP012
HWDA	Healing Abdomen wound; 70&90 min post incision	pCMVSport 3.0	LP012
HWEA	Healing Abdomen Wound; 15 days post incision	pCMVSport 3.0	LP012
HWJA	Healing Abdomen Wound;21&29 days	pCMVSport 3.0	LP012
HNAL	Human Tongue, frac 2	pSport1	LP012
НМЈА	H. Meniingima, M6	pSport1	LP012
HMKA HMKB HMKC HMKD HMKE	H. Meningima, M1	pSport1	LP012
НОГА	Ovarian Tumor I, OV5232	pSport1	LP012
HCFA HCFB HCFC HCFD	T-Cell PHA 16 hrs	pSport1	LP012
HCFL HCFM HCFN HCFO	T-Cell PHA 24 hrs	pSport1	LP012
НММА НММВ НММС	Spleen metastic melanoma	pSport1	LP012
HTDA	Human Tonsil, Lib 3	pSport1	LP012
HDBA	Human Fetal Thymus	pSport1	LP012
HDUA	Pericardium	pSport1	LP012
HBZA	Prostate, BPH, Lib 2	pSport1	LP012
HWCA	Larynx tumor	pSport1	LP012
HWKA	Normal lung	pSport1	LP012
HSMB	Bone marrow stroma,treated	pSport1	LP012

Normal trachea		- у		
HBHM Normal trachea, pSport1 LP012 HLFC Human Larynx pSport1 LP012 HLRB Siebben Polyposis pSport1 LP012 HNIA Mammary Gland pSport1 LP012 HNIB Palate carcinoma pSport1 LP012 HNIB Palate carcinoma pSport1 LP012 HNKA Palate normal pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HMZA Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pancreas normal PCA4 No pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HUCA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HUKE HUMAN Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random primed Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human umbilical Vein Endothelial cells, Lambda ZAP II LP013 cells, fract. B Human Umbilical Vein Endothelial cells, Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013	Libraries owned by Catalog	Catalog Description	Vector	ATCC
HLFC Human Larynx pSport1 LP012 HLRB Siebben Polyposis pSport1 LP012 HNIA Mammary Gland pSport1 LP012 HNIB Palate carcinoma pSport1 LP012 HNIB Palate carcinoma pSport1 LP012 HNKA Palate normal pSport1 LP012 HMKA Palate normal pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HMZA Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HWAA Pancreas normal PCA4 No pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HUKA HUKA HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HUKA HUKA HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HTDA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 HMEB Human Umbilical Vein Endothelial cells, Lambda ZAP II LP013 HCDC HLQD Hepatocellular tumor, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, differentially expressed HSUS H. Striatum Depression, subtracted pBluescript LP013				Deposit
HLRB Siebben Polyposis pSport1 LP012 HNIA Mammary Gland pSport1 LP012 HNIB Palate carcinoma pSport1 LP012 HNKA Palate normal pSport1 LP012 HNKA Palate normal pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HMZA Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HUKA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HUKE HFFA Human Fetal Brain, random primed Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 Fract. B HUSH Human Umbilical Vein Endothelial cells, Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTPS Human Whole 6 week Old Embryo (II), pBluescript LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013			pSport1	LP012
HNIA Mammary Gland pSport1 LP012 HNJB Palate carcinoma pSport1 LP012 HNKA Palate normal pSport1 LP012 HNKA Palate normal pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HUCA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HHFA Human Fetal Brain, random primed Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 Fract. B HUSH Human Umbilical Vein Endothelial cells, Lambda ZAP II LP013 HLQC HLQD Hepatocellular tumor, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Whole 6 week Old Embryo (II), pBluescript LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013			pSport1	LP012
HNJB Palate carcinoma pSport1 LP012 HNKA Palate normal pSport1 LP012 HNKA Palate normal pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HABG Cheek Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HICA Tongue carcinoma pSport1 LP012 HICA Tongue carcinoma pSport1 LP013 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human umbilical Vein Endothelial cells, Lambda ZAP II LP013 fract. B HUSH Human Umbilical Vein Endothelial cells, fact. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Whole 6 week Old Embryo (II), pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, subtracted pBluescript LP013 HSUS Supt cells, cyclohexamide treated, differentially expressed HSUS HSDS H. Striatum Depression, subtracted pBluescript LP013			pSport l	LP012
HNKA Palate normal pSport1 LP012 HMZA Pharynx carcinoma pSport1 LP012 HABG Cheek Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HUKA HUKA HUKA HUKA HUKA HUMA Uterine Cancer Lambda ZAP II LP013 HTVAA HUMA PART PART PART PART PART PART PART PAR		Mammary Gland	pSport1	LP012
HMZA Pharynx carcinoma pSport1 LP012 HABG Cheek Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HWAA Pancreas normal PCA4 No pSport1 LP012 HIVAA Pancreas normal PCA4 No pSport1 LP012 HICA Tongue carcinoma pSport1 LP013 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random primed Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 fract. B HUSH Human Umbilical Vein Endothelial cells, Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Whole 6 week Old Embryo (II), pBluescript LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS H. Striatum Depression, subtracted pBluescript LP013		Palate carcinoma	pSport1	LP012
HABG Cheek Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HICA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HTFA Human Fetal Brain, random primed Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 fract. B HUSH Human Umbilical Vein Endothelial cells, Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Whole 6 week Old Embryo (II), pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLIS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS HSDS H. Striatum Depression, subtracted pBluescript LP013	HNKA	Palate normal	pSport1	LP012
HMZM Pharynx Carcinoma pSport1 LP012 HMZM Pharynx Carcinoma pSport1 LP012 HDRM Larynx Carcinoma pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HICA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HTUKE HUMAN Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 fract. B HUSH Human Umbilical Vein Endothelial Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 subt HHPS Human Hippocampus, subtracted pBluescript LP013 HL1S LNCAP, differential expression pBluescript LP013 HL1S LNCAP, differential expression pBluescript LP013 HL1S Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS H. Striatum Depression, subtracted pBluescript LP013	HMZA	Pharynx carcinoma	pSport1	LP012
HDRM Larynx Carcinoma pSport1 LP012 HVAA Pancreas normal PCA4 No pSport1 LP012 HICA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HTVAA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 Fract. B HUSH Human Umbilical Vein Endothelial Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 subt HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013	HABG	Cheek Carcinoma	pSport1	LP012
HVAA Pancreas normal PCA4 No pSport1 LP012 HICA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HUKE HIFFA Human Fetal Brain, random primed Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random primed Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, fract. B HUSH Human Umbilical Vein Endothelial Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HF6S Human Whole 6 week Old Embryo (II), pBluescript LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUS Supt cells, cyclohexamide treated, differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013	HMZM	Pharynx Carcinoma	pSport1	LP012
HICA Tongue carcinoma pSport1 LP012 HUKA HUKB HUKC HUKD Human Uterine Cancer Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HUSH Human Umbilical Vein Endothelial cells, fract. B HUSH Human Umbilical Vein Endothelial cells, fract. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Whole 6 week Old Embryo (II), subt HUMAN Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, gBluescript LP013 HSUS Supt cells, cyclohexamide treated, differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013	HDRM	Larynx Carcinoma	pSport1	LP012
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HUKE HFFA Human Fetal Brain, random primed Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri HBQA Early Stage Human Brain, random primed Human microvascular Endothelial cells, Lambda ZAP II LP013 HUSH Human Umbilical Vein Endothelial cells, fract. A, re-excision HLQC HLQD Hepatocellular tumor, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HUMAN Whole 6 week Old Embryo (II), pBluescript LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 differentially expressed HSUS H. Striatum Depression, subtracted pBluescript LP013	HICA	Tongue carcinoma	pSport1	LP012
HFFA Human Fetal Brain, random primed Lambda ZAP II LP013 HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 HUSH Human Umbilical Vein Endothelial Lambda ZAP II LP013 cells, fract. B Lambda ZAP II LP013 cells, fract. A, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HF6S Human Whole 6 week Old Embryo (II), pBluescript LP013 HHPS Human Hippocampus, subtracted pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLHS HLHT Early Stage Human Lung, Subtracted pBluescript LP013 HSUS Supt cells, cyclohexamide treated, subtracted pBluescript LP013 HSUT Supt cells, cyclohexamide treated, differentially expressed HSUT Bluescript LP013 HSDS H. Striatum Depression, subtracted pBluescript LP013	HUKA HUKB HUKC HUKD	Human Uterine Cancer	Lambda ZAP II	LP013
HTUA Activated T-cell labeled with 4-thioluri Lambda ZAP II LP013 HBQA Early Stage Human Brain, random primed Lambda ZAP II LP013 HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 HUSH Human Umbilical Vein Endothelial Lambda ZAP II LP013 HLQC HLQD Hepatocellular tumor, re-excision Lambda ZAP II LP013 HTWJ HTWK HTWL Resting T-cell, re-excision Lambda ZAP II LP013 HHPS Human Whole 6 week Old Embryo (II), pBluescript LP013 HLIS LNCAP, differential expression pBluescript LP013 HLHS HLHT Early Stage Human Lung, Subtracted pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 HSUT Supt cells, cyclohexamide treated, differentially expressed HSUS H. Striatum Depression, subtracted pBluescript LP013	HUKE			
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HMEB Human microvascular Endothelial cells, Lambda ZAP II LP013 fract. B HUSH Human Umbilical Vein Endothelial Lambda ZAP II LP013 cells, fract. A, re-excision HLQC HLQD Hepatocellular tumor, re-excision HTWJ HTWK HTWL Resting T-cell, re-excision Human Whole 6 week Old Embryo (II), pBluescript LP013 HHPS Human Hippocampus, subtracted HUSH	HTUA	Activated T-cell labeled with 4-thioluri	Lambda ZAP II	LP013
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HHPS Human Hippocampus, subtracted pBluescript LP013 HL1S LNCAP, differential expression pBluescript LP013 HLHS HLHT Early Stage Human Lung, Subtracted pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 subtracted PBluescript LP013 HSUT Supt cells, cyclohexamide treated, pBluescript LP013 differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013	HTWJ HTWK HTWL	Resting T-cell, re-excision	Lambda ZAP II	LP013
subt HHPS Human Hippocampus, subtracted pBluescript LP013 HL18 LNCAP, differential expression pBluescript LP013 HLHS HLHT Early Stage Human Lung, Subtracted pBluescript LP013 HSUS Supt cells, cyclohexamide treated, pBluescript LP013 subtracted HSUT Supt cells, cyclohexamide treated, pBluescript LP013 differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013	HF6S	Human Whole 6 week Old Embryo (II),	pBluescript	LP013
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HSUT Supt cells, cyclohexamide treated, pBluescript LP013 differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013	HLHS HLHT	Early Stage Human Lung, Subtracted	pBluescript	LP013
HSUT Supt cells, cyclohexamide treated, pBluescript LP013 differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013	HSUS		<u> </u>	LP013
differentially expressed HSDS H. Striatum Depression, subtracted pBluescript LP013		subtracted		
HSDS H. Striatum Depression, subtracted pBluescript LP013	HSUT	Supt cells, cyclohexamide treated,	pBluescript	LP013
ung-z		differentially expressed		
IID77	HSDS	H. Striatum Depression, subtracted	pBluescript	LP013
Human Pituitary, Subtracted VII pBluescript LP013	HPTZ	Human Pituitary, Subtracted VII	pBluescript	LP013
HSDX H. Striatum Depression, subt II pBluescript LP013	HSDX	H. Striatum Depression, subt II	· ·	
HSDZ H. Striatum Depression, subt pBluescript LP013	HSDZ	H. Striatum Depression, subt	•	
HPBA HPBB HPBC HPBD HPBE Human Pineal Gland pBluescript SK- LP013	ІРВА НРВВ НРВС НРВО НРВІ	E Human Pineal Gland	<u></u>	
HRTA Colorectal Tumor pBluescript SK- LP013	IRTA	Colorectal Tumor		

Libraries owned by Catalog	Catalog Description	Vector	ATCC
			Deposit
HSBA HSBB HSBC HSBM	HSC172 cells	pBluescript SK-	LP013
HJAA HJAB HJAC HJAD	Jurkat T-cell G1 phase	pBluescript SK-	LP013
НЈВА НЈВВ НЈВС НЈВD	Jurkat T-cell, S1 phase	pBluescript SK-	LP013
HTNA HTNB	Human Thyroid	pBluescript SK-	LP013
НАНА НАНВ	Human Adult Heart	Uni-ZAP XR	LP013'
HE6A	Whole 6 week Old Embryo	Uni-ZAP XR	LP013
HFCA HFCB HFCC HFCD HFCE	Human Fetal Brain	Uni-ZAP XR	LP013
HFKC HFKD HFKE HFKF	Human Fetal Kidney	Uni-ZAP XR	LP013
HFKG	·		
HGBA HGBD HGBE HGBF	Human Gall Bladder	Uni-ZAP XR	LP013
HGBG			
HPRA HPRB HPRC HPRD	Human Prostate	Uni-ZAP XR	LP013
HTEA HTEB HTEC HTED HTEE		Uni-ZAP XR	LP013
HTTA HTTB HTTC HTTD HTTE	Human Testes Tumor	Uni-ZAP XR	LP013
НҮВА НҮВВ	Human Fetal Bone	Uni-ZAP XR	LP013
HFLA	Human Fetal Liver	Uni-ZAP XR	LP013
ННГВ ННГС ННГО ННГЕ ННГГ	Human Fetal Heart	Uni-ZAP XR	LP013
HUVB HUVC HUVD HUVE	Human Umbilical Vein, End. remake	Uni-ZAP XR	LP013
HTHB HTHC HTHD	Human Thymus	Uni-ZAP XR	LP013
HSTA HSTB HSTC HSTD	Human Skin Tumor	Uni-ZAP XR	LP013
HTAA HTAB HTAC HTAD	Human Activated T-cells	Uni-ZAP XR	LP013
HTAE			
HFEA HFEB HFEC	Human Fetal Epithelium (skin)	Uni-ZAP XR	LP013
НЈРА НЈРВ НЈРС НЈРД	Human Jurkat Membrane Bound	Uni-ZAP XR	LP013
	Polysomes		
HESA	Human Epithelioid Sarcoma	Uni-ZAP XR	LP013
HALS	Human Adult Liver, Subtracted	Uni-ZAP XR	LP013
HFTA HFTB HFTC HFTD	Human Fetal Dura Mater	Uni-ZAP XR	LP013
НСАА НСАВ НСАС	Cem cells, cyclohexamide treated	Uni-ZAP XR	LP013 .
HRGA HRGB HRGC HRGD	Raji Cells, cyclohexamide treated	Uni-ZAP XR	LP013
НЕ9А НЕ9В НЕ9С НЕ9D НЕ9E	Nine Week Old Early Stage Human	Uni-ZAP XR	LP013
HSFA	Human Fibrosarcoma	Uni-ZAP XR	LP013
HATA HATB HATC HATD	Human Adrenal Gland Tumor	Uni-ZAP XR	LP013
HATE			
HTRA	Human Trachea Tumor	Uni-ZAP XR	LP013
HE2A HE2D HE2E HE2H HE2I	12 Week Old Early Stage Human	Uni-ZAP XR	LP013
HE2B HE2C HE2F HE2G HE2P	12 Week Old Early Stage Human, II	Uni-ZAP XR	LP013
INEA HNEB HNEC HNED	Human Neutrophil	Uni-ZAP XR	LP013
INEE	-		

Libraries owned by Catalog	Catalog Description	Vector	ATCC
	Catalog Description	vector ,	
HBGA	Human Primary Breast Cancer	Uni-ZAP XR	Deposit
HPTS HPTT HPTU	Human Pituitary, subtracted	Uni-ZAP XR	LP013
HMQA HMQB HMQC HMQD	Human Activated Monocytes		LP013
HOAA HOAB HOAC	Human Osteosarcoma	Uni-ZAP XR	LP013
HTOA HTOD HTOE HTOF		Uni-ZAP XR	LP013
HTOG	human tonsils	Uni-ZAP XR	, LP013
HMGB	Human OB MG63 control fraction I		
HOPB		Uni-ZAP XR	LP013 .
	Human OB HOS control fraction I	Uni-ZAP XR	LP013
НООВ	Human OB HOS treated (1 nM E2) fraction I	Uni-ZAP XR	LP013
HAUA HAUB HAUC	Amniotic Cells - TNF induced	Uni-ZAP XR	LP013
HAQA HAQB HAQC HAQD	Amniotic Cells - Primary Culture	Uni-ZAP XR	LP013
HROA HROC	HUMAN STOMACH	Uni-ZAP XR	LP013
НВЈА НВЈВ НВЈС НВЈО НВЈЕ	HUMAN B CELL LYMPHOMA	Uni-ZAP XR	LP013
HODA HODB HODC HODD	human ovarian cancer	Uni-ZAP XR	LP013
НСРА	Corpus Callosum	Uni-ZAP XR	LP013
HSOA	stomach cancer (human)	Uni-ZAP XR	LP013
HERA	SKIN	Uni-ZAP XR	LP013
HMDA	Brain-medulloblastoma	Uni-ZAP XR	LP013
HGLA HGLB HGLD	Glioblastoma	Uni-ZAP XR	LP013
HWTA HWTB HWTC	wilm's tumor	Uni-ZAP XR	LP013
НЕАА	H. Atrophic Endometrium	Uni-ZAP XR	LP013
НАР N НАРО НАРР НАРQ НАРR	Human Adult Pulmonary;re-excision	Uni-ZAP XR	LP013
HLTG HLTH	Human T-cell lymphoma;re-excision	Uni-ZAP XR	LP013
HAHC HAHD HAHE	Human Adult Heart;re-excision	Uni-ZAP XR	LP013
HAGA HAGB HAGC HAGD HAGE	Human Amygdala	Uni-ZAP XR	LP013
НЅЈА НЅЈВ НЅЈС	Smooth muscle-ILb induced	Uni-ZAP XR	LP013
НЅНА НЅНВ НЅНС	Smooth muscle, IL1b induced	Uni-ZAP XR	LP013
HPWA HPWB HPWC HPWD HPWE	Prostate BPH	Uni-ZAP XR	LP013
НРІА НРІВ НРІС	LNCAP prostate cell line	Uni-ZAP XR	LP013
НРЈА НРЈВ НРЈС	PC3 Prostate cell line	Uni-ZAP XR	LP013
НВТА	Bone Marrow Stroma, TNF&LPS ind	Uni-ZAP XR	LP013
НМСГ НМСС НМСН НМСІ	Macrophage-oxLDL; re-excision	Uni-ZAP XR	LP013
НМСЈ			
HAGG HAGH HAGI	Human Amygdala;re-excision	Uni-ZAP XR	LP013
HACA	H. Adipose Tissue	Uni-ZAP XR	LP013

Libraries owned by Catalog	Catalog Description	Vector	ATCC
HIVED			Deposit
HKFB	K562 + PMA (36 hrs), re-excision	ZAP Express	LP013
HCWT HCWU HCWV	CD34 positive cells (cord blood),re-ex	ZAP Express	LP013
HBWA	Whole brain	ZAP Express	LP013
HBXA HBXB HBXC HBXD	Human Whole Brain #2 - Oligo dT > 1.5Kb	ZAP Express	LP013
HAVM	Temporal cortex-Alzheizmer	pT-Adv	LP014
HAVT	Hippocampus, Alzheimer Subtracted	pT-Adv	LP014
HHAS	CHME Cell Line	Uni-ZAP XR	LP014
HAJR	Larynx normal	pSport 1	LP014
HWLE HWLF HWLG HWLH	Colon Normal	pSport 1	LP014
HCRM HCRN HCRO	Colon Carcinoma	pSport I	LP014
HWLI HWLJ HWLK	Colon Normal	pSport 1	LP014
HWLQ HWLR HWLS HWLT	Colon Tumor	pSport 1	LP014
НВГМ	Gastrocnemius Muscle	pSport 1	LP014
HBOD HBOE	Quadriceps Muscle	pSport 1	LP014
НВКО НВКЕ	Soleus Muscle	pSport 1	LP014
HCCM	Pancreatic Langerhans	pSport 1	LP014
HWGA	Larynx carcinoma	pSport I	LP014
HWGM HWGN	Larynx carcinoma	pSport 1	LP014
HWLA HWLB HWLC	Normal colon	pSport 1	LP014
IWLM HWLN	Colon Tumor	pSport I	LP014
IVAM HVAN HVAO	Pancreas Tumor	pSport 1	LP014
HWGQ	Larynx carcinoma	pSport I	LP014
AAQM HAQN	Salivary Gland	pSport 1	LP014
IASM	Stomach; normal	pSport I	LP014
IBCM	Uterus; normal	pSport 1	LP014
ICDM	Testis; normal	pSport 1	LP014
IDJM	Brain; normal	pSport I	LP014
IEFM	Adrenal Gland, normal	pSport I	LP014
IBAA	Rectum normal	pSport I	LP014
IFDM	Rectum tumour	pSport 1	LP014
IGAM	Colon, normal	pSport I	LP014
НММ	Colon, tumour	pSport 1	LP014
CLB HCLC	Human Lung Cancer	Lambda Zap II	LP015
RLA	L1 Cell line	ZAP Express	LP015
HAM	Hypothalamus, Alzheimer's	pCMVSport 3.0	LP015
KBA	Ku 812F Basophils Line	pSport 1	LP015
S2S	Saos2, Dexamethosome Treated	pSport 1	LP016

Libraries owned by Catalog	Catalog Description	Vector	ATCC
			Deposit
HASA	Lung Carcinoma A549 TNFalpha	pSport 1	LP016
	activated		
HTFM	TF-1 Cell Line GM-CSF Treated	pSport 1	LP016
HYAS	Thyroid Tumour	pSport 1	LP016
HUTS	Larynx Normal	pSport 1	LP016
НХОА	Larynx Tumor	pSport 1	LP016
НЕАН	Ea.hy.926 cell line	pSport 1	LP016
HINA	Adenocarcinoma Human	pSport 1	LP016
HRMA	Lung Mesothelium	pSport 1	LP016
HLCL	Human Pre-Differentiated Adipocytes	Uni-Zap XR	LP017
HS2A	Saos2 Cells	pSport 1	LP020
HS21	Saos2 Cells; Vitamin D3 Treated	pSport I	LP020
HUCM	CHME Cell Line, untreated	pSport 1	LP020
HEPN	Aryepiglottis Normal	pSport 1	LP020
HPSN	Sinus Piniformis Tumour	pSport 1	LP020
HNSA	Stomach Normal	pSport I	LP020
HNSM	Stomach Tumour	pSport 1	LP020
HNLA	Liver Normal Met5No	pSport 1	LP020
HUTA	Liver Tumour Met 5 Tu	pSport 1	LP020
HOCN	Colon Normal	pSport 1	LP020
НОСТ	Colon Tumor	pSport 1	LP020
HTNT	Tongue Tumour	pSport 1	LP020
HLXN	Larynx Normal	pSport 1	LP020
HLXT	Larynx Tumour	pSport 1	LP020
HTYN	Thymus	pSport 1	LP020
HPLN	Placenta	pSport 1	LP020
HTNG	Tongue Normal	pSport 1	LP020
HZAA	Thyroid Normal (SDCA2 No)	pSport 1	LP020
HWES	Thyroid Thyroiditis	pSport 1	LP020
HFHD	Ficolled Human Stromal Cells, 5Fu	pTrip1Ex2	LP021
•	treated		
HFHM,HFHN	Ficolled Human Stromal Cells,	pTrip1Ex2	LP021
	Untreated		
HPCI	Hep G2 Cells, lambda library	lambda Zap-CMV	LP021
		XR	
нвса,нвсв,нвсс	H. Lymph node breast Cancer	Uni-ZAP XR	LP021
НСОК	Chondrocytes	pSPORTI	LP022
HDCA, HDCB, HDCC	Dendritic Cells From CD34 Cells	pSPORT1	LP022

Libraries owned by Catalog	Catalog Description	Vector	ATCC ·
,			Deposit
HDMA, HDMB	CD40 activated monocyte dendritic	pSPORT1	LP022
,	cells		
HDDM, HDDN, HDDO	LPS activated derived dendritic cells	pSPORT1	LP022
HPCR	Hep G2 Cells, PCR library	lambda Zap-CMV	LP022
(XR	
HAAA, HAAB, HAAC	Lung, Cancer (4005313A3): Invasive	pSPORT1	LP022
4	Poorly Differentiated Lung		,
	Adenocarcinoma		
HIPA, HIPB, HIPC	Lung, Cancer (4005163 B7): Invasive,	pSPORT1	LP022
•	Poorly Diff. Adenocarcinoma,		
	Metastatic		
ноон, нооі	Ovary, Cancer: (4004562 B6) Papillary	pSPORT1	LP022
	Serous Cystic Neoplasm, Low		
	Malignant Pot	1	
HIDA	Lung, Normal: (4005313 B1)	pSPORT1	LP022
HUJA,HUJB,HUJC,HUJD,HUJE	B-Cells	pCMVSport 3.0	LP022
HNOA,HNOB,HNOC,HNOD	Ovary, Normal: (9805C040R)	pSPORTI	LP022
HNLM	Lung, Normal: (4005313 B1)	pSPORTI	LP022
HSCL	Stromal Cells	pSPORTI	LP022
HAAX	Lung, Cancer: (4005313 A3) Invasive	pSPORT1	LP022
	Poorly-differentiated Metastatic lung		
	adenocarcinoma		
HUUA,HUUB,HUUC,HUUD	B-cells (unstimulated)	pTrip1Ex2	LP022
HWWA,HWWB,HWWC,HWWD,	B-cells (stimulated)	pSPORTI	LP022
HWWE,HWWF,HWWG			
HCCC	Colon, Cancer: (9808C064R)	pCMVSport 3.0	LP023
HPDO HPDP HPDQ HPDR HPD	Ovary, Cancer (9809C332): Poorly	pSport 1	LP023
	differentiated adenocarcinoma		ľ
НРСО НРСР НРСО НРСТ	Ovary, Cancer (15395A1F): Grade II	pSport 1	LP023
•	Papillary Carcinoma		
НОСМ НОСО НОСР НОСО	Ovary, Cancer: (15799A1F) Poorly	pSport 1	LP023
	differentiated carcinoma		
НСВМ НСВО НСВО	Breast, Cancer: (4004943 A5)	pSport 1	LP023
INBT HNBU HNBV	Breast, Normal: (4005522B2)	pSport I	LP023
НВСР НВСQ	Breast, Cancer: (4005522 A2)	pSport 1	LP023
HBCJ	Breast, Cancer: (9806C012R)	pSport 1	LP023
ISAM HSAN	Stromal cells 3.88	pSport I	LP023
IVCA HVCB HVCC HVCD	Ovary, Cancer: (4004332 A2)	pSport 1	LP023
	•	• •	1

Libraries owned by Catalog	Catalog Description	Vector	ATCC
			Deposit
HSCP HSCQ	stromal cell clone 2.5	pSport 1	LP023
HUXA	Breast Cancer: (4005385 A2)	pSport 1	LP023
HCOM HCON HCOO HCOP HCOQ	Ovary, Cancer (4004650 A3): Well- Differentiated Micropapillary Serous Carcinoma	pSport 1	LP023
HBNM	Breast, Cancer: (9802C020E)	pSport 1	LP023
HVVA HVVB HVVC HVVD HVVE	Human Bone Marrow, treated	pSport 1	LP023

[0888] Two nonlimiting examples are provided below for isolating a particular clone from the deposited sample of plasmid cDNAs cited for that clone in Table 7. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to the nucleotide sequence of SEQ ID NO:X.

Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The oligonucleotide is labeled, for instance, with ³²P-γ-ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).) The plasmid mixture is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above. The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening (e.g., Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1.93 to 1.104), or other techniques known to those of skill in the art.

[0890] Alternatively, two primers of 17-20 nucleotides derived from both ends of the nucleotide sequence of SEQ ID NO:X are synthesized and used to amplify the desired cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 μl of reaction mixture with 0.5 ug of the above cDNA template. A convenient reaction mixture is

1.5-5 mM MgCl₂, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

- [0891] Several methods are available for the identification of the 5' or 3' non-coding portions of a gene which may not be present in the deposited clone. These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols which are well known in the art. For instance, a method similar to 5' RACE is available for generating the missing 5' end of a desired full-length transcript. (Fromont-Racine et al., Nucleic Acids Res. 21(7):1683-1684 (1993).)
- [0892] Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be sequenced and used to generate the full length gene.
- although poly-A+ RNA can be used. The RNA preparation can then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged RNA which may interfere with the later RNA ligase step. The phosphatase should then be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase.
- [0894] This modified RNA preparation is used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the

gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the desired gene.

Example 2: Isolation of Genomic Clones Corresponding to a Polynucleotide

[0895] A human genomic P1 library (Genomic Systems, Inc.) is screened by PCR using primers selected for the sequence corresponding to SEQ ID NO:X according to the method described in Example 1. (See also, Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Edn., (1989), Cold Spring Harbor Laboratory Press).

Example 3: Tissue specific expression analysis

[0896] The Human Genome Sciences, Inc. (HGS) database is derived from sequencing tissue and/or disease specific cDNA libraries. Libraries generated from a particular tissue are selected and the specific tissue expression pattern of EST groups or assembled contigs within these libraries is determined by comparison of the expression patterns of those groups or contigs within the entire database. ESTs and assembled contigs which show tissue specific expression are selected.

in the case of an assembled contig, the clone from which the 5' most EST sequence was generated, is obtained from the catalogued library of clones and the insert amplified by PCR using methods known in the art. The PCR product is denatured and then transferred in 96 or 384 well format to a nylon membrane (Schleicher and Scheull) generating an array filter of tissue specific clones. Housekeeping genes, maize genes, and known tissue specific genes are included on the filters. These targets can be used in signal normalization and to validate assay sensitivity. Additional targets are included to monitor probe length and specificity of hybridization.

[0898] Radioactively labeled hybridization probes are generated by first strand cDNA synthesis per the manufacturer's instructions (Life Technologies) from mRNA/RNA samples prepared from the specific tissue being analyzed (e.g.,

reproductive system, cancers of the reproductive system, or, more specifically, prostate, prostate cancer, ovarian, ovarian cancer, etc.). The hybridization probes are purified by gel exclusion chromatography, quantitated, and hybridized with the array filters in hybridization bottles at 65°C overnight. The filters are washed under stringent conditions and signals are captured using a Fuji phosphorimager.

[0899] Data is extracted using AIS software and following background subtraction, signal normalization is performed. This includes a normalization of filterwide expression levels between different experimental runs. Genes that are differentially expressed in the tissue of interest are identified.

Example 4: Chromosomal Mapping of the Polynucleotides

[0900] An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This primer set is then used in a polymerase chain reaction under the following set of conditions: 30 seconds, 95°C; 1 minute, 56°C; 1 minute, 70°C. This cycle is repeated 32 times followed by one 5 minute cycle at 70°C. Human, mouse, and hamster DNA is used as template in addition to a somatic cell hybrid panel containing individual chromosomes or chromosome fragments (Bios, Inc). The reactions are analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

Example 5: Bacterial Expression of a Polypeptide

[0901] A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as BamHI and XbaI, at the 5' end of the primers in order to clone the amplified

product into the expression vector. For example, BamHI and XbaI correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (Amp^r), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a ribosome binding site (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

The pQE-9 vector is digested with BamHI and Xbal and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, which expresses the lacI repressor and also confers kanamycin resistance (Kan^r). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis.

[0903] Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D. 600) of between 0.4 and 0.6. IPTG (Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1 mM. IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased gene expression.

[0904] Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4°C. The cell debris is removed by centrifugation, and the supernatant containing the polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from QIAGEN, Inc., supra). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., supra).

[0905] Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8. The column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then

washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

[0906] The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins are eluted by the addition of 250 mM immidazole. Immidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

In addition to the above expression vector, the present invention further includes an expression vector, called pHE4a (ATCC Accession Number 209645, deposited on February 25, 1998) which contains phage operator and promoter elements operatively linked to a polynucleotide of the present invention. This vector contains: 1) a neomycinphosphotransferase gene as a selection marker, 2) an E. coli origin of replication, 3) a T5 phage promoter sequence, 4) two lac operator sequences, 5) a Shine-Delgarno sequence, and 6) the lactose operon repressor gene (lacIq). The origin of replication (oriC) is derived from pUC19 (LTI, Gaithersburg, MD). The promoter and operator sequences are made synthetically.

[0908] DNA can be inserted into the pHE4a by restricting the vector with Ndel and Xbal, BamHI, XhoI, or Asp718, running the restricted product on a gel, and isolating the larger fragment (the stuffer fragment should be about 310 base pairs). The DNA insert is generated according to the PCR protocol described in Example 1, using PCR primers having restriction sites for NdeI (5' primer) and XbaI, BamHI, XhoI, or Asp718 (3' primer). The PCR insert is gel purified and restricted with compatible enzymes. The insert and vector are ligated according to standard protocols.

[0909] The engineered vector could easily be substituted in the above protocol to express protein in a bacterial system.

Example 6: Purification of a Polypeptide from an Inclusion Body

[0910] The following alternative method can be used to purify a polypeptide expressed in E coli when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

- [0911] Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.
- [0912] The cells are then lysed by passing the solution through a microfluidizer (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.
- [0913] The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the pellet is discarded and the polypeptide containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.
- [0914] Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.
- [0915] To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 μm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH

6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0. Both column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A₂₈₀ monitoring of the effluent. Fractions containing the polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

[0917] The resultant polypeptide should exhibit greater than 95% purity after the above refolding and purification steps. No major contaminant bands should be observed from Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded. The purified protein can also be tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

Example 7: Cloning and Expression of a Polypeptide in a Baculovirus Expression System

[0918] In this example, the plasmid shuttle vector pA2 is used to insert a polynucleotide into a baculovirus to express a polypeptide. This expression vector contains the strong polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites such as BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from *E. coli* under control of a weak Drosophila promoter in the same orientation, followed by the polyadenylation signal

of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

- [0919] Many other baculovirus vectors can be used in place of the vector above, such as pAc373, pVL941, and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription, translation, secretion and the like, including a signal peptide and an inframe AUG as required. Such vectors are described, for instance, in Luckow et al., Virology 170:31-39 (1989).
- [0920] Specifically, the cDNA sequence contained in the deposited clone, including the AUG initiation codon, is amplified using the PCR protocol described in Example 1. If a naturally occurring signal sequence is used to produce the polypeptide of the present invention, the pA2 vector does not need a second signal peptide. Alternatively, the vector can be modified (pA2 GP) to include a baculovirus leader sequence, using the standard methods described in Summers et al., "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures," Texas Agricultural Experimental Station Bulletin No. 1555 (1987).
- [0921] The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.
- [0922] The plasmid is digested with the corresponding restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("Geneclean" BIO 101 Inc., La Jolla, Ca.).
- [0923] The fragment and the dephosphorylated plasmid are ligated together with T4 DNA ligase. E. coli HB101 or other suitable E. coli hosts such as XL-1 Blue (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria containing the plasmid are identified by digesting DNA from individual colonies and analyzing the digestion product by gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA sequencing.

Five μg of a plasmid containing the polynucleotide is co-transfected with 1.0 μg of a commercially available linearized baculovirus DNA ("BaculoGoldTM baculovirus DNA", Pharmingen, San Diego, CA), using the lipofection method described by Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One μg of BaculoGoldTM virus DNA and 5 μg of the plasmid are mixed in a sterile well of a microtiter plate containing 50 μl of serum-free Grace's medium (Life Technologies Inc., Gaithersburg, MD). Afterwards, 10 μl Lipofectin plus 90 μl Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27° C. The transfection solution is then removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27° C for four days.

performed, as described by Summers and Smith, *supra*. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200 μl of Grace's medium and the suspension containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

[0926] To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5 μCi of ³⁵S-methionine and 5 μCi ³⁵S-cysteine (available from Amersham) are added. The cells

are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

[0927] Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the produced protein.

Example 8: Expression of a Polypeptide in Mammalian Cells

The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which mediates the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLVI, HIVI and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

[0929] Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden), pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSport 2.0, and pCMVSport 3.0. Mammalian host cells that could be used include, human Hela, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

[0930] Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transfection with a selectable marker such as DHFR, gpt, neomycin, or hygromycin allows the identification and isolation of the transfected cells.

The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing cell lines that carry several hundred or even several thousand copies of the gene of interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta, 1097:107-143 (1990); Page, M. J. and Sydenham, M. A., Biotechnology 9:64-68 (1991).) Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., Biochem J. 227:277-279 (1991); Bebbington et al., Bio/Technology 10:169-175 (1992). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

Derivatives of the plasmid pSV2-dhfr (ATCC Accession No. 37146), the expression vectors pC4 (ATCC Accession No. 209646) and pC6 (ATCC Accession No.209647) contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., Molecular and Cellular Biology, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., Cell 41:521-530 (1985).) Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, XbaI and Asp718, facilitate the cloning of the gene of interest. The vectors also contain the 3' intron, the polyadenylation and termination signal of the rat preproinsulin gene, and the mouse DHFR gene under control of the SV40 early promoter.

[0933] Specifically, the plasmid pC6, for example, is digested with appropriate restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from a 1% agarose gel.

[0934] A polynucleotide of the present invention is amplified according to the protocol outlined in Example 1. If a naturally occurring signal sequence is used to produce the polypeptide of the present invention, the vector does not need a second signal peptide. Alternatively, if a naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., International Publication No. WO 96/34891.)

[0935] The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment

then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

[0936] The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. E. coli HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene is used for [0937] transfection. Five µg of the expression plasmid pC6 or pC4 is cotransfected with 0.5 μg of the plasmid pSVneo using lipofectin (Felgner et al., supra). The plasmid pSV2neo contains a dominant selectable marker, the neo gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μ M, 2 μ M, 5 μ M, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 - 200 µM. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

Example 9: Protein Fusions

[0938] The polypeptides of the present invention are preferably fused to other proteins. These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG domains, and maltose binding protein facilitates purification. (See Example 5; see also EP A 394,827; Traunecker, et al., Nature 331:84-86 (1988).) Similarly, fusion to IgG-1, IgG-3, and albumin increases the halflife time *in vivo*. Nuclear localization signals

fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or homodimers can increase or decrease the activity of a fusion protein. Fusion proteins can also create chimeric molecules having more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by modifying the following protocol, which outlines the fusion of a polypeptide to an IgG molecule, or the protocol described in Example 5.

- [0939] Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate cloning into an expression vector, preferably a mammalian expression vector.
- [0940] For example, if pC4 (ATCC Accession No. 209646) is used, the human Fc portion can be ligated into the BamHI cloning site. Note that the 3' BamHI site should be destroyed. Next, the vector containing the human Fc portion is re-restricted with BamHI, linearizing the vector, and a polynucleotide of the present invention, isolated by the PCR protocol described in Example 1, is ligated into this BamHI site. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.
- [0941] If the naturally occurring signal sequence is used to produce the polypeptide of the present invention, pC4 does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., International Publication No. WO 96/34891.)

[0942] Human IgG Fc region:

GGGATCCGGAGCCCAAATCTTCTGACAAAACTCACACATGCCCACCGTGC
CCAGCACCTGAATTCGAGGGTGCACCGTCAGTCTTCCTCTTCCCCCCAAAA
CCCAAGGACACCCTCATGATCTCCCGGACTCCTGAGGTCACATGCGTGGTG
GTGGACGTAAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGA
CGGCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTAC
AACAGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGACTGG

CTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCAAC
CCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCAC
AGGTGTACACCCTGCCCCCATCCCGGGATGAGCTGACCAAGAACCAGGTC
AGCCTGACCTGCCTGGTCAAAGGCTTCTATCCAAGCGACATCGCCGTGGAG
TGGGAGAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGT
GCTGGACTCCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAA
GAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGG
CTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAAT
GAGTGCGACGGCCGCGACTCTAGAGGAT (SEQ ID NO: 1)

Example 10: Production of an Antibody from a Polypeptide

Hybridoma Technology

[0943] The antibodies of the present invention can be prepared by a variety of methods. (See, Current Protocols, Chapter 2.) As one example of such methods, cells expressing polypeptide of the present invention are administered to an animal to induce the production of sera containing polyclonal antibodies. In a preferred method, a preparation of polypeptide of the present invention is prepared and purified to render it substantially free of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

Monoclonal antibodies specific for polypeptide of the present invention are prepared using hybridoma technology (Kohler et al., Nature 256:495 (1975); Kohler et al., Eur. J. Immunol. 6:511 (1976); Kohler et al., Eur. J. Immunol. 6:292 (1976); Hammerling et al., in: Monoclonal Antibodies and T-Cell Hybridomas, Elsevier, N.Y., pp. 563-681 (1981)). In general, an animal (preferably a mouse) is immunized with polypeptide of the present invention or, more preferably, with a secreted polypeptide of the present invention-expressing cell. Such polypeptide-expressing cells are cultured in any suitable tissue culture medium, preferably in Earle's modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at about 56°C), and supplemented with about 10 g/l of nonessential amino acids, about 1,000 U/ml of penicillin, and about 100 μg/ml of streptomycin.

The splenocytes of such mice are extracted and fused with a suitable myeloma cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line (SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are selectively maintained in HAT medium, and then cloned by limiting dilution as described by Wands et al. (Gastroenterology 80:225-232 (1981)). The hybridoma cells obtained through such a selection are then assayed to identify clones which secrete antibodies capable of binding the polypeptide of the present invention.

Alternatively, additional antibodies capable of binding to polypeptide of the present invention can be produced in a two-step procedure using anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the polypeptide of the present invention-specific antibody can be blocked by polypeptide of the present invention. Such antibodies comprise anti-idiotypic antibodies to the polypeptide of the present invention-specific antibody and are used to immunize an animal to induce formation of further polypeptide of the present invention-specific antibodies.

antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric and humanized antibodies are known in the art and are discussed herein. (See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., International Publication No. WO 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 (1985).)

Isolation Of Antibody Fragments Directed Against Polypeptide of the Present Invention From A Library Of scFvs

[0948] Naturally occurring V-genes isolated from human PBLs are constructed into a library of antibody fragments which contain reactivities against polypeptide of the present invention to which the donor may or may not have been exposed (see e.g., U.S. Patent 5,885,793 incorporated herein by reference in its entirety).

Rescue of the Library. A library of scFvs is constructed from the RNA of human PBLs as described in International Publication No. WO 92/01047. To rescue phage displaying antibody fragments, approximately 10° E. coli harboring the phagemid are used to inoculate 50 ml of 2xTY containing 1% glucose and 100 μg/ml of ampicillin (2xTY-AMP-GLU) and grown to an O.D. of 0.8 with shaking. Five ml of this culture is used to inoculate 50 ml of 2xTY-AMP-GLU, 2 x 108 TU of delta gene 3 helper (M13 delta gene III, see International Publication No. WO 92/01047) are added and the culture incubated at 37°C for 45 minutes without shaking and then at 37°C for 45 minutes with shaking. The culture is centrifuged at 4000 r.p.m. for 10 min. and the pellet resuspended in 2 liters of 2xTY containing 100 μg/ml ampicillin and 50 ug/ml kanamycin and grown overnight. Phage are prepared as described in International Application No. WO 92/01047.

does not encode gene III is prepared as follows: M13 delta gene III helper phage does not encode gene III protein, hence the phage(mid) displaying antibody fragments have a greater avidity of binding to antigen. Infectious M13 delta gene III particles are made by growing the helper phage in cells harboring a pUC19 derivative supplying the wild type gene III protein during phage morphogenesis. The culture is incubated for 1 hour at 37° C without shaking and then for a further hour at 37°C with shaking. Cells are spun down (IEC-Centra 8,400 r.p.m. for 10 min), resuspended in 300 ml 2xTY broth containing 100 μg ampicillin/ml and 25 μg kanamycin/ml (2xTY-AMP-KAN) and grown overnight, shaking at 37°C. Phage particles are purified and concentrated from the culture medium by two PEG-precipitations (Sambrook et al., 1990), resuspended in 2 ml PBS and passed through a 0.45 μm filter (Minisart NML; Sartorius) to give a final concentration of approximately 10¹³ transducing units/ml (ampicillin-resistant clones).

[0951] Panning of the Library. Immunotubes (Nunc) are coated overnight in PBS with 4 ml of either 100 μg/ml or 10 μg/ml of a polypeptide of the present invention. Tubes are blocked with 2% Marvel-PBS for 2 hours at 37°C and then washed 3 times

in PBS. Approximately 10¹³ TU of phage is applied to the tube and incubated for 30 minutes at room temperature tumbling on an over and under turntable and then left to stand for another 1.5 hours. Tubes are washed 10 times with PBS 0.1% Tween-20 and 10 times with PBS. Phage are eluted by adding 1 ml of 100 mM triethylamine and fotating 15 minutes on an under and over turntable after which the solution is immediately neutralized with 0.5 ml of 1.0M Tris-HCl, pH 7.4. Phage are then used to infect 10 ml of mid-log E. coli TG1 by incubating eluted phage with bacteria for 30 minutes at 37°C. The E. coli are then plated on TYE plates containing 1% glucose and 100 µg/ml ampicillin. The resulting bacterial library is then rescued with delta gene 3 helper phage as described above to prepare phage for a subsequent round of selection. This process is then repeated for a total of 4 rounds of affinity purification with tubewashing increased to 20 times with PBS, 0.1% Tween-20 and 20 times with PBS for rounds 3 and 4.

Characterization of Binders. Eluted phage from the 3rd and 4th rounds of selection are used to infect E. coli HB 2151 and soluble scFv is produced (Marks, et al., 1991) from single colonies for assay. ELISAs are performed with microtitre plates coated with 10 pg/ml of the polypeptide of the present invention in 50 mM bicarbonate pH 9.6. Clones positive in ELISA are further characterized by PCR fingerprinting (see, e.g., International Application No. WO 92/01047) and then by sequencing. These ELISA positive clones may also be further characterized by techniques known in the art, such as, for example, epitope mapping, binding affinity, receptor signal transduction, ability to block or competitively inhibit antibody/antigen binding, and competitive agonistic or antagonistic activity.

Example 11: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

[0953] RNA isolated from entire families or individual patients presenting with a phenotype of interest (such as a disease) is isolated. cDNA is then generated from these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in

SEQ ID NO:X; and/or the nucleotide sequence of the cDNA contained in Clone ID NO:Z. Suggested PCR conditions consist of 35 cycles at 95 degrees C for 30 seconds; 60-120 seconds at 52-58 degrees C; and 60-120 seconds at 70 degrees C, using buffer solutions described in Sidransky et al., Science 252:706 (1991).

[0954] PCR products are then sequenced using primers labeled at their 5' end with T4 polynucleotide kinase, employing SequiTherm Polymerase (Epicentre Technologies). The intron-exon boundaries of selected exons is also determined and genomic PCR products analyzed to confirm the results. PCR products harboring suspected mutations are then cloned and sequenced to validate the results of the direct sequencing.

[0955] PCR products are cloned into T-tailed vectors as described in Holton et al., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals are identified by mutations not present in unaffected individuals.

[0956] Genomic rearrangements are also observed as a method of determining alterations in a gene corresponding to a polynucleotide. Genomic clones isolated according to Example 2 are nick-translated with digoxigenindeoxy-uridine 5'-triphosphate (Boehringer Manheim), and FISH performed as described in Johnson et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is carried out using a vast excess of human cot-1 DNA for specific hybridization to the corresponding genomic locus.

[0957] Chromosomes are counterstained with 4,6-diamino-2-phenylidole and propidium iodide, producing a combination of C- and R-bands. Aligned images for precise mapping are obtained using a triple-band filter set (Chroma Technology, Brattleboro, VT) in combination with a cooled charge-coupled device camera (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson et al., Genet. Anal. Tech. Appl., 8:75 (1991).) Image collection, analysis and chromosomal fractional length measurements are performed using the ISee Graphical Program System. (Inovision Corporation, Durham, NC.) Chromosome alterations of the genomic region hybridized by the probe are identified as insertions, deletions, and translocations. These alterations are used as a diagnostic marker for an associated disease.

Example 12: Method of Detecting Abnormal Levels of a Polypeptide in a Biological Sample

[0958] A polypeptide of the present invention can be detected in a biological sample, and if an increased or decreased level of the polypeptide is detected, this polypeptide is a marker for a particular phenotype. Methods of detection are numerous, and thus, it is understood that one skilled in the art can modify the following assay to fit their particular needs.

[0959] For example, antibody-sandwich ELISAs are used to detect polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method described in Example 10. The wells are blocked so that non-specific binding of the polypeptide to the well is reduced.

[0960] The coated wells are then incubated for > 2 hours at RT with a sample containing the polypeptide. Preferably, serial dilutions of the sample should be used to validate results. The plates are then washed three times with deionized or distilled water to remove unbound polypeptide.

[0961] Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at a concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove unbound conjugate.

[0962] Add 75 ul of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl phosphate (NPP) substrate solution to each well and incubate 1 hour at room temperature. Measure the reaction by a microtiter plate reader. Prepare a standard curve, using serial dilutions of a control sample, and plot polypeptide concentration on the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale). Interpolate the concentration of the polypeptide in the sample using the standard curve.

Example 13: Formulation

[0963] The invention also provides methods of treatment and/or prevention of diseases or disorders (such as, for example, any one or more of the diseases or disorders disclosed herein) by administration to a subject of an effective amount of a Therapeutic. By "Therapeutic" is meant polynucleotides or polypeptides of the invention (including fragments, analogs, derivatives and variants thereof), agonists or antagonists thereof, and/or antibodies thereto (including fragments thereof), in combination with a pharmaceutically acceptable carrier type (e.g., a sterile carrier).

[0964] The Therapeutic will be formulated and dosed in a fashion consistent with good medical practice, taking into account the clinical condition of the individual patient (especially the side effects of treatment with the Therapeutic alone), the site of delivery, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" for purposes herein is thus determined by such considerations.

As a general proposition, the total pharmaceutically effective amount of the Therapeutic administered parenterally per dose will be in the range of about lug/kg/day to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If given continuously, the Therapeutic is typically administered at a dose rate of about 1 ug/kg/hour to about 50 ug/kg/hour, either by 1-4 injections per day or by continuous subcutaneous infusions, for example, using a mini-pump. An intravenous bag solution may also be employed. The length of treatment needed to observe changes and the interval following treatment for responses to occur appears to vary depending on the desired effect.

[0966] Therapeutics can be are administered orally, rectally, parenterally, intracistemally, intravaginally, intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), bucally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any. The term "parenteral" as used herein refers to modes of administration which include

intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

Therapeutics of the invention are also suitably administered by sustained-release systems. Suitable examples of sustained-release Therapeutics are administered orally, rectally, parenterally, intracistemally, intravaginally, intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), bucally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers to modes of administration which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

[0968] Therapeutics of the invention are also suitably administered by sustained-release systems. Suitable examples of sustained-release Therapeutics include suitable polymeric materials (such as, for example, semi-permeable polymer matrices in the form of shaped articles, e.g., films, or mirocapsules), suitable hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, and sparingly soluble derivatives (such as, for example, a sparingly soluble salt).

[0969] Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman et al., Biopolymers 22:547-556 (1983)), poly (2- hydroxyethyl methacrylate) (Langer et al., J. Biomed. Mater. Res. 15:167-277 (1981), and Langer, Chem. Tech. 12:98-105 (1982)), ethylene vinyl acetate (Langer et al., Id.) or poly-D- (-)-3-hydroxybutyric acid (EP 133,988).

Therapeutics of the invention (see generally, Langer, Science 249:1527-1533 (1990); Treat et al., in Liposomes in the Therapy of Infectious Disease and Cancer, Lopez-Berestein and Fidler (eds.), Liss, New York, pp. 317 -327 and 353-365 (1989)). Liposomes containing the Therapeutic are prepared by methods known per se: DE 3,218,121; Epstein et al., Proc. Natl. Acad. Sci. (USA) 82:3688-3692 (1985); Hwang et al., Proc. Natl. Acad. Sci. (USA) 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes are of the small

(about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal Therapeutic.

[0971] In yet an additional embodiment, the Therapeutics of the invention are delivered by way of a pump (see Langer, supra; Sefton, CRC Crit. Ref. Biomed. Eng. 14:201 (1987); Buchwald et al., Surgery 88:507 (1980); Saudek et al., N. Engl. J. Med. 321:574 (1989)).

[0972] Other controlled release systems are discussed in the review by Langer (Science 249:1527-1533 (1990)).

[0973] For parenteral administration, in one embodiment, the Therapeutic is formulated generally by mixing it at the desired degree of purity, in a unit dosage injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations employed and is compatible with other ingredients of the formulation. For example, the formulation preferably does not include oxidizing agents and other compounds that are known to be deleterious to the Therapeutic.

[0974] Generally, the formulations are prepared by contacting the Therapeutic uniformly and intimately with liquid carriers or finely divided solid carriers or both. Then, if necessary, the product is shaped into the desired formulation. Preferably the carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood of the recipient. Examples of such carrier vehicles include water, saline, Ringer's solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl oleate are also useful herein, as well as liposomes.

that enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at the dosages and concentrations employed, and include buffers such as phosphate, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose or its

derivatives, glucose, manose, or dextrins; chelating agents such as EDTA; sugar alcohols such as mannitol or sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates, poloxamers, or PEG.

- [0976] The Therapeutic is typically formulated in such vehicles at a concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of polypeptide salts.
- [0977] Any pharmaceutical used for therapeutic administration can be sterile. Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutics generally are placed into a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.
- [0978] Therapeutics ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials are filled with 5 ml of sterile-filtered 1% (w/v) aqueous Therapeutic solution, and the resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the lyophilized Therapeutic using bacteriostatic Water-for-Injection.
- [0979] The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the Therapeutics of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the Therapeutics may be employed in conjunction with other therapeutic compounds.
- [0980] The Therapeutics of the invention may be administered alone or in combination with adjuvants. Adjuvants that may be administered with the Therapeutics of the invention include, but are not limited to, alum, alum plus deoxycholate (ImmunoAg), MTP-PE (Biocine Corp.), QS21 (Genentech, Inc.), BCG, and MPL. In a specific embodiment, Therapeutics of the invention are administered in combination with alum. In another specific embodiment, Therapeutics of the invention are administered in combination with QS-21. Further adjuvants that may be

administered with the Therapeutics of the invention include, but are not limited to, Monophosphoryl lipid immunomodulator, AdjuVax 100a, QS-21, QS-18, CRL1005, Aluminum salts, MF-59, and Virosomal adjuvant technology. Vaccines that may be administered with the Therapeutics of the invention include, but are not limited to, vaccines directed toward protection against MMR (measles, mumps, rubella), polio, varicella, tetanus/diptheria, hepatitis A, hepatitis B, haemophilus influenzae B, whooping cough, pneumonia, influenza, Lyme's Disease, rotavirus, cholera, yellow fever, Japanese encephalitis, poliomyelitis, rabies, typhoid fever, and pertussis. Combinations may be administered either concomitantly, e.g., as an admixture, separately but simultaneously or concurrently; or sequentially. presentations in which the combined agents are administered together as a therapeutic mixture, and also procedures in which the combined agents are administered separately but simultaneously, e.g., as through separate intravenous lines into the same individual. Administration "in combination" further includes the separate administration of one of the compounds or agents given first, followed by the second.

[0981] The Therapeutics of the invention may be administered alone or in combination with other therapeutic agents. Therapeutic agents that may be administered in combination with the Therapeutics of the invention, include but not limited to, other members of the TNF family, chemotherapeutic agents, antibiotics, steroidal and non-steroidal anti-inflammatories, conventional immunotherapeutic agents, cytokines and/or growth factors. Combinations may be administered either concomitantly, e.g., as an admixture, separately but simultaneously or concurrently; or sequentially. This includes presentations in which the combined agents are administered together as a therapeutic mixture, and also procedures in which the combined agents are administered separately but simultaneously, e.g., as through separate intravenous lines into the same individual. Administration "in combination" further includes the separate administration of one of the compounds or agents given first, followed by the second.

[0982] In one embodiment, the Therapeutics of the invention are administered in combination with members of the TNF family. TNF, TNF-related or TNF-like molecules that may be administered with the Therapeutics of the invention include, but are not limited to, soluble forms of TNF-alpha, lymphotoxin-alpha (LT-alpha, also

known as TNF-beta), LT-beta (found in complex heterotrimer LT-alpha2-beta), OPGL, FasL, CD27L, CD30L, CD40L, 4-1BBL, DcR3, OX40L, TNF-gamma (International Publication No. WO 96/14328), AIM-I (International Publication No. WO 97/33899), endokine-alpha (International Publication No. WO 98/07880), TR6 (International Publication No. WO 98/30694), OPG, and neutrokine-alpha (International Publication No. WO 98/18921, OX40, and nerve growth factor (NGF), and soluble forms of Fas, CD30, CD27, CD40 and 4-IBB, TR2 (International Publication No. WO 96/34095), DR3 (International Publication No. WO 97/33904), DR4 (International Publication No. WO 98/30693), TR6 (International Publication No. WO 98/30694), TR7 (International Publication No. WO 98/56892),TR10 (International Publication No. WO 98/54202), 312C2 (International Publication No. WO 98/06842), and TR12, and soluble forms CD154, CD70, and CD153.

[0983] In certain embodiments, Therapeutics of the invention are administered in combination with antiretroviral agents, nucleoside reverse transcriptase inhibitors, non-nucleoside reverse transcriptase inhibitors, and/or protease inhibitors. Nucleoside reverse transcriptase inhibitors that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, RETROVIR TM (zidovudine/AZT), VIDEX™ (didanosine/ddI), HIVID™ (zalcitabine/ddC), ZERIT™ (stavudine/d4T), **EPIVIR™** (lamivudine/3TC), and COMBIVIR™ (zidovudine/lamivudine). Non-nucleoside reverse transcriptase inhibitors that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, VIRAMUNE™ (nevirapine), RESCRIPTOR™ (delavirdine), and SUSTIVA™ (efavirenz). Protease inhibitors that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, CRIXIVANTM (indinavir), NORVIR™ (ritonavir), INVIRASE™ (saquinavir), and VIRACEPT™ (nelfinavir). In a specific embodiment, antiretroviral agents, nucleoside reverse transcriptase inhibitors, non-nucleoside reverse transcriptase inhibitors, and/or protease inhibitors may be used in any combination with Therapeutics of the invention to treat AIDS and/or to prevent or treat HIV infection.

In other embodiments, Therapeutics of the invention may be administered [0984] in combination with anti-opportunistic infection agents. Anti-opportunistic agents that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, TRIMETHOPRIM-SULFAMETHOXAZOLE™, DAPSONE™, PENTAMIDINE™, ATOVAQUONE™, ISONIAZID™, RIFAMPIN™, PYRAZINAMIDE™, ETHAMBUTOL™, RIFABUTIN™, CLARITHROMYCIN™, AZITHROMYCIN™, GANCICLOVIR™, FOSCARNET™, CIDOFOVIR™, FLUCONAZOLE™, ITRACONAZOLE™, KETOCONAZOLE™, ACYCLOVIR™, FAMCICOLVIR™, PYRIMETHAMINE™, LEUCOVORIN™, NEUPOGEN™ (filgrastim/G-CSF), and LEUKINE™ (sargramostim/GM-CSF). In a specific embodiment, Therapeutics of the invention are used in any combination with TRIMETHOPRIM-SULFAMETHOXAZOLE™, DAPSONE™, PENTAMIDINE™, and/or ATOVAQUONE™ to prophylactically treat or prevent an opportunistic Pneumocystis carinii pneumonia infection. In another specific embodiment, Therapeutics of the invention are used in any combination with ISONIAZID™, RIFAMPIN™, PYRAZINAMIDE™, and/or ETHAMBUTOL™ to prophylactically treat or prevent an opportunistic Mycobacterium avium complex infection. In another specific embodiment, Therapeutics of the invention are used in any combination with RIFABUTIN™, CLARITHROMYCIN™, and/or AZITHROMYCIN™ to prophylactically treat or prevent an opportunistic Mycobacterium tuberculosis infection. In another specific embodiment, Therapeutics of the invention are used in any combination with GANCICLOVIR™, FOSCARNET™, and/or CIDOFOVIR™ to prophylactically treat or prevent an opportunistic cytomegalovirus infection. In another specific embodiment, Therapeutics of the invention are used in any combination with FLUCONAZOLE™, ITRACONAZOLE™, and/or KETOCONAZOLE™ to prophylactically treat or prevent an opportunistic fungal infection. In another specific embodiment, Therapeutics of the invention are used in any combination with ACYCLOVIR™ and/or $FAMCICOLVIR^{TM}$ to prophylactically treat or prevent an opportunistic herpes simplex virus type I and/or type II infection. In another specific embodiment, Therapeutics of the invention are used in any combination

PYRIMETHAMINE™ and/or LEUCOVORIN™ to prophylactically treat or prevent an opportunistic *Toxoplasma gondii* infection. In another specific embodiment, Therapeutics of the invention are used in any combination with LEUCOVORIN™ and/or NEUPOGEN™ to prophylactically treat or prevent an opportunistic bacterial infection.

- [0985] In a further embodiment, the Therapeutics of the invention are administered in combination with an antiviral agent. Antiviral agents that may be administered with the Therapeutics of the invention include, but are not limited to, acyclovir, ribavirin, amantadine, and remantidine.
- [0986] In a further embodiment, the Therapeutics of the invention arc administered in combination with an antibiotic agent. Antibiotic agents that may be administered with the Therapeutics of the invention include, but are not limited to, amoxicillin, beta-lactamases, aminoglycosides, beta-lactam (glycopeptide), beta-lactamases, Clindamycin, chloramphenicol, cephalosporins, ciprofloxacin, ciprofloxacin, erythromycin, fluoroquinolones, macrolides, metronidazole, penicillins, quinolones, rifampin, streptomycin, sulfonamide, tetracyclines, trimethoprim, trimethoprim-sulfamthoxazole, and vancomycin.
- [0987] 'Conventional nonspecific immunosuppressive agents, that may be administered in combination with the Therapeutics of the invention include, but are not limited to, steroids, cyclosporine, cyclosporine analogs, cyclophosphamide methylprednisone, prednisone, azathioprine, FK-506, 15-deoxyspergualin, and other immunosuppressive agents that act by suppressing the function of responding T cells.
- 109881 In specific embodiments, Therapeutics of the invention are administered in combination with immunosuppressants. Immunosuppressants preparations that may be administered with the Therapeutics of the invention include, but are not limited to, ORTHOCLONE™ (OKT3), SANDIMMUNE™/NEORAL™/SANGDYA™ (cyclosporin), PROGRAF™ (tacrolimus). CELLCEPT™ (mycophenolate), Azathioprine, glucorticosteroids, and RAPAMUNE™ (sirolimus). In a specific embodiment, immunosuppressants may be used to prevent rejection of organ or bone marrow transplantation.

In an additional embodiment, Therapeutics of the invention are administered alone or in combination with one or more intravenous immune globulin preparations. Intravenous immune globulin preparations that may be administered with the Therapeutics of the invention include, but not limited to, GAMMARTM, IVEEGAMTM, SANDOGLOBULINTM, GAMMAGARD S/DTM, and GAMIMUNETM. In a specific embodiment, Therapeutics of the invention are administered in combination with intravenous immune globulin preparations in transplantation therapy (e.g., bone marrow transplant).

[0990] In an additional embodiment, the Therapeutics of the invention are administered alone or in combination with an anti-inflammatory agent. Anti-inflammatory agents that may be administered with the Therapeutics of the invention include, but are not limited to, glucocorticoids and the nonsteroidal anti-inflammatories, aminoarylcarboxylic acid derivatives, arylacetic acid derivatives, arylbutyric acid derivatives, arylcarboxylic acids, arylpropionic acid derivatives, pyrazoles, pyrazolones, salicylic acid derivatives, thiazinecarboxamides, e-acetamidocaproic acid, S-adenosylmethionine, 3-amino-4-hydroxybutyric acid, amixetrine, bendazac, benzydamine, bucolome, difenpiramide, ditazol, emorfazone, guaiazulene, nabumetone, nimesulide, orgotein, oxaceprol, paranyline, perisoxal, pifoxime, proquazone, proxazole, and tenidap.

[0991] In an additional embodiment, the compositions of the invention are administered alone or in combination with an anti-angiogenic agent. Anti-angiogenic agents that may be administered with the compositions of the invention include, but are not limited to, Angiostatin (Entremed, Rockville, MD), Troponin-1 (Boston Life Sciences, Boston, MA), anti-Invasive Factor, retinoic acid and derivatives thereof, paclitaxel (Taxol), Suramin, Tissue Inhibitor of Metalloproteinase-1, Tissue Inhibitor of Metalloproteinase-2, VEGI, Plasminogen Activator Inhibitor-1, Plasminogen Activator Inhibitor-2, and various forms of the lighter "d group" transition metals.

[0992] Lighter "d group" transition metals include, for example, vanadium, molybdenum, tungsten, titanium, niobium, and tantalum species. Such transition metal species may form transition metal complexes. Suitable complexes of the above-mentioned transition metal species include oxo transition metal complexes.

[0993] Representative examples of vanadium complexes include oxo vanadium complexes such as vanadate and vanadyl complexes. Suitable vanadate complexes include metavanadate and orthovanadate complexes such as, for example, ammonium metavanadate, sodium metavanadate, and sodium orthovanadate. Suitable vanadyl complexes include, for example, vanadyl acetylacetonate and vanadyl sulfate including vanadyl sulfate hydrates such as vanadyl sulfate mono- and trihydrates.

include oxo complexes. Suitable oxo tungsten and molybdenum complexes also include oxo complexes. Suitable oxo tungsten complexes include tungstate and tungsten oxide complexes. Suitable tungstate complexes include ammonium tungstate, calcium tungstate, sodium tungstate dihydrate, and tungstic acid. Suitable tungsten oxides include tungsten (IV) oxide and tungsten (VI) oxide. Suitable oxo molybdenum complexes include molybdate, molybdenum oxide, and molybdenyl complexes. Suitable molybdate complexes include ammonium molybdate and its hydrates, sodium molybdate and its hydrates, and potassium molybdate and its hydrates. Suitable molybdenum oxides include molybdenum (VI) oxide, molybdenum (VI) oxide, and molybdic acid. Suitable molybdenyl complexes include, for example, molybdenyl acetylacetonate. Other suitable tungsten and molybdenum complexes include hydroxo derivatives derived from, for example, glycerol, tartaric acid, and sugars.

[0995] A wide variety of other anti-angiogenic factors may also be utilized within the context of the present invention. Representative examples include, but are not limited to, platelet factor 4; protamine sulphate; sulphated chitin derivatives (prepared from queen crab shells), (Murata et al., Cancer Res. 51:22-26 (1991)); Sulphated Polysaccharide Peptidoglycan Complex (SP-PG) (the function of this compound may be enhanced by the presence of steroids such as estrogen, and tamoxifen citrate); Staurosporine; modulators of matrix metabolism, including for example, proline analogs, cishydroxyproline, d,L-3,4-dehydroproline, Thiaproline, alpha, alphadipyridyl, aminopropionitrile fumarate; 4-propyl-5-(4-pyridinyl)-2(3H)-oxazolone; Methotrexate; Mitoxantrone; Heparin; Interferons; 2 Macroglobulin-serum; ChIMP-3 (Pavloff et al., J. Bio. Chem. 267:17321-17326, (1992)); Chymostatin (Tomkinson et al., Biochem J. 286:475-480 (1992)); Cyclodextrin Tetradecasulfate; Eponemycin; Camptothecin; Fumagillin (Ingber et al., Nature 348:555-557, 1990); Gold Sodium

Thiomalate ("GST"; Matsubara and Ziff, J. Clin. Invest. 79:1440-1446, 1987); anticollagenase-serum; alpha2-antiplasmin (Holmes et al., J. Biol. Chem. 262(4):1659-1664 (1987)); Bisantrene (National Cancer Institute); Lobenzarit disodium (N-(2)-carboxyphenyl-4- chloroanthronilic acid disodium or "CCA"; (Takeuchi et al., Agents Actions 36:312-316, 1992); and metalloproteinase inhibitors such as BB94.

Additional anti-angiogenic factors that may also be utilized within the context of the present invention include Thalidomide, (Celgene, Warren, NJ); Angiostatic steroid; AGM-1470 (H. Brem and J. Folkman J Pediatr. Surg. 28:445-51 (1993)); an integrin alpha v beta 3 antagonist (C. Storgard et al., J Clin. Invest. 103:47-54 (1999)); carboxynaminolmidazole; Carboxyamidotriazole (CAI) (National Cancer Institute, Bethesda, MD); Conbretastatin A-4 (CA4P) (OXiGENE, Boston, MA); Squalamine (Magainin Pharmaceuticals, Plymouth Meeting, PA); TNP-470, (Tap Pharmaceuticals, Deerfield, IL); ZD-0101 AstraZeneca (London, UK); APRA (CT2584); Benefin, Byrostatin-1 (SC339555); CGP-41251 (PKC 412); CM101; Dexrazoxane (ICRF187); DMXAA; Endostatin; Flavopridiol; Genestein; GTE; ImmTher; Iressa (ZD1839); Octreotide (Somatostatin); Panretin; Penacillamine; Photopoint; PI-88; Prinomastat (AG-3340) Purlytin; Suradista (FCE26644); Tamoxifen (Nolvadex); Tazarotene; Tetrathiomolybdate; Xeloda (Capecitabine); and 5-Fluorouracil.

Anti-angiogenic agents that may be administed in combination with the compounds of the invention may work through a variety of mechanisms including, but not limited to, inhibiting proteolysis of the extracellular matrix, blocking the function of endothelial cell-extracellular matrix adhesion molecules, by antagonizing the function of angiogenesis inducers such as growth factors, and inhibiting integrin receptors expressed on proliferating endothelial cells. Examples of anti-angiogenic inhibitors that interfere with extracellular matrix proteolysis and which may be administered in combination with the compositons of the invention include, but are not lmited to, AG-3340 (Agouron, La Jolla, CA), BAY-12-9566 (Bayer, West Haven, CT), BMS-275291 (Bristol Myers Squibb, Princeton, NJ), CGS-27032A (Novartis, East Hanover, NJ), Marimastat (British Biotech, Oxford, UK), and Metastat (Aeterna, St-Foy, Quebec). Examples of anti-angiogenic inhibitors that act by blocking the function of endothelial cell-extracellular matrix adhesion molecules and which may be

administered in combination with the compositons of the invention include, but are not lmited to, EMD-121974 (Merck KcgaA Darmstadt, Germany) and Vitaxin (Ixsys, La Jolla, CA/Medimmune, Gaithersburg, MD). Examples of anti-angiogenic agents that act by directly antagonizing or inhibiting angiogenesis inducers and which may be administered in combination with the compositons of the invention include, but are not lmited to, Angiozyme (Ribozyme, Boulder, CO), Anti-VEGF antibody (Genentech, S. San Francisco, CA), PTK-787/ZK-225846 (Novartis, Basel, Switzerland), SU-101 (Sugen, S. San Francisco, CA), SU-5416 (Sugen/ Pharmacia Upjohn, Bridgewater, NJ), and SU-6668 (Sugen). Other anti-angiogenic agents act to indirectly inhibit angiogenesis. Examples of indirect inhibitors of angiogenesis which may be administered in combination with the compositons of the invention include, but are not limited to, IM-862 (Cytran, Kirkland, WA), Interferon-alpha, IL-12 (Roche, Nutley, NJ), and Pentosan polysulfate (Georgetown University, Washington, DC).

[0998] In particular embodiments, the use of compositions of the invention in combination with anti-angiogenic agents is contemplated for the treatment, prevention, and/or amelioration of an autoimmune disease, such as for example, an autoimmune disease described herein.

[0999] In a particular embodiment, the use of compositions of the invention in combination with anti-angiogenic agents is contemplated for the treatment, prevention, and/or amelioration of arthritis. In a more particular embodiment, the use of compositions of the invention in combination with anti-angiogenic agents is contemplated for the treatment, prevention, and/or amelioration of rheumatoid arthritis.

[01000] In another embodiment, the polynucleotides encoding a polypeptide of the present invention are administered in combination with an angiogenic protein, or polynucleotides encoding an angiogenic protein. Examples of angiogenic proteins that may be administered with the compositions of the invention include, but are not limited to, acidic and basic fibroblast growth factors, VEGF-1, VEGF-2, VEGF-3, epidermal growth factor alpha and beta, platelet-derived endothelial cell growth factor, platelet-derived growth factor, tumor necrosis factor alpha, hepatocyte growth factor, insulin like growth factor, colony stimulating factor, macrophage colony stimulating factor, granulocyte/macrophage colony stimulating factor, and nitric oxide synthase.

[01001] In another embodiment, compostions of the invention are administered in combination with a chemotherapeutic agent. Chemotherapeutic agents that may be administered with the compositions of the invention include, but are not limited to, antibiotic derivatives (e.g., doxorubicin, bleomycin, daunorubicin, and dactinomycin); antiestrogens tamoxifen); antimetabolites (e.g., (e.g., fluorouracil. floxuridine, interferon alpha-2b, glutamic acid, plicamycin, methotrexate. mercaptopurine, and 6-thioguanine); cytotoxic agents (e.g., carmustine, BCNU, CCNU, cytosine arabinoside, cyclophosphamide, hydroxyurea, procarbazine, mitomycin, busulfan, cis-platin, and vincristine sulfate); hormones (e.g., medroxyprogesterone, estramustine phosphate sodium, ethinyl estradiol, megestrol acetate, methyltestosterone, estradiol, diethylstilbestrol diphosphate, chlorotrianisene, and testolactone); nitrogen mustard derivatives (e.g., mephalen, chorambucil, mechlorethamine (nitrogen mustard) and thiotepa); steroids and combinations (e.g., bethamethasone sodium phosphate); and others (e.g., dicarbazine, asparaginase, mitotane, vincristine sulfate, vinblastine sulfate, and etoposide).

[01002] In a specific embodiment, Therapeutics of the invention are administered in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, and prednisone) or any combination of the components of CHOP. In another embodiment, Therapeutics of the invention are administered in combination with Rituximab. In a further embodiment, Therapeutics of the invention are administered with Rituxmab and CHOP, or Rituxmab and any combination of the components of CHOP.

In an additional embodiment, the Therapeutics of the invention are administered in combination with cytokines. Cytokines that may be administered with the Therapeutics of the invention include, but are not limited to, IL2, IL3, IL4, IL5, IL6, IL7, IL10, IL12, IL13, IL15, anti-CD40, CD40L, IFN-gamma and TNF-alpha. In another embodiment, Therapeutics of the invention may be administered with any interleukin, including, but not limited to, IL-1alpha, IL-1beta, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-14, IL-15, IL-16, IL-17, IL-18, IL-19, IL-20, and IL-21.

[01004] In an additional embodiment, the Therapeutics of the invention are administered in combination with angiogenic proteins. Angiogenic proteins that may

be administered with the Therapeutics of the invention include, but are not limited to, Glioma Derived Growth Factor (GDGF), as disclosed in European Patent Number EP-399816; Platelet Derived Growth Factor-A (PDGF-A), as disclosed in European Patent Number EP-682110; Platelet Derived Growth Factor-B (PDGF-B), as disclosed in European Patent Number EP-282317; Placental Growth Factor (PIGF), as disclosed in International Publication Number WO 92/06194; Placental Growth Factor-2 (PIGF-2). as disclosed in Hauser et al., Gorwth Factors, 4:259-268 (1993); Vascular Endothelial Growth Factor (VEGF), as disclosed in International Publication Number WO 90/13649; Vascular Endothelial Growth Factor-A (VEGF-A), as disclosed in European Patent Number EP-506477; Vascular Endothelial Growth Factor-2 (VEGF-2), as disclosed in International Publication Number WO 96/39515; Vascular Endothelial Growth Factor B (VEGF-3); Vascular Endothelial Growth Factor B-186 (VEGF-B186), as disclosed in International Publication Number WO 96/26736; Vascular Endothelial Growth Factor-D (VEGF-D), as disclosed in International Publication Number WO 98/02543; Vascular Endothelial Growth Factor-D (VEGF-D), as disclosed in International Publication Number WO 98/07832; and Vascular Endothelial Growth Factor-E (VEGF-E), as disclosed in German Patent Number DE19639601. The above mentioned references are herein incorporated by reference in their entireties.

- In an additional embodiment, the Therapeutics of the invention are administered in combination with hematopoietic growth factors. Hematopoietic growth factors that may be administered with the Therapeutics of the invention include, but are not limited to, LEUKINETM (SARGRAMOSTIMTM) and NEUPOGENTM (FILGRASTIMTM).
- [01006] In an additional embodiment, the Therapeutics of the invention are administered in combination with Fibroblast Growth Factors. Fibroblast Growth Factors that may be administered with the Therapeutics of the invention include, but are not limited to, FGF-1, FGF-2, FGF-3, FGF-4, FGF-5, FGF-6, FGF-7, FGF-8, FGF-9, FGF-10, FGF-11, FGF-12, FGF-13, FGF-14, and FGF-15.
- [01007] In additional embodiments, the Therapeutics of the invention are administered in combination with other therapeutic or prophylactic regimens, such as, for example, radiation therapy.

Example 14: Method of Treating Decreased Levels of the Polypeptide

of an increased level of a polypeptide of the invention in the body comprising administering to such an individual a composition comprising a therapeutically effective amount of an agonist of the invention (including polypeptides of the invention). Moreover, it will be appreciated that conditions caused by a decrease in the standard or normal expression level of a polypeptide of the present invention in an individual can be treated by administering the agonist or antagonist of the present invention. Thus, the invention also provides a method of treatment of an individual in need of an increased level of the polypeptide comprising administering to such an individual a Therapeutic comprising an amount of the agonist or antagonist to increase the activity level of the polypeptide in such an individual.

[01009] For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the agonist or antagonist for six consecutive days. The exact details of the dosing scheme, based on administration and formulation, are provided in Example 13.

Example 15: Method of Treating Increased Levels of the Polypeptide

[01010] The present invention also relates to a method of treating an individual in need of a decreased level of a polypeptide of the invention in the body comprising administering to such an individual a composition comprising a therapeutically effective amount of an antagonist of the invention (including polypeptides and antibodies of the invention).

[01011] In one example, antisense technology is used to inhibit production of a polypeptide of the present invention. This technology is one example of a method of decreasing levels of a polypeptide, due to a variety of etiologies, such as cancer.

[01012] For example, a patient diagnosed with abnormally increased levels of a polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5,

2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The formulation of the antisense polynucleotide is provided in Example 13.

Example 16: Method of Treatment Using Gene Therapy-Ex Vivo

[01013] One method of gene therapy transplants fibroblasts, which are capable of expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and separated into small pieces. Small chunks of the tissue are placed on a wet surface of a tissue culture flask, approximately ten pieces are placed in each flask. The flask is turned upside down, closed tight and left at room temperature over night. After 24 hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to the bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS, penicillin and streptomycin) is added. The flasks are then incubated at 37 degree C for approximately one week.

[01014] At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The monolayer is trypsinized and scaled into larger flasks.

[01015] pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long terminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and HindIII and, subsequently treated with calf intestinal phosphatase. The linear vector is fractionated on agarose gel and purified, using glass beads.

[01016] The cDNA encoding a polypeptide of the present invention can be amplified using PCR primers which correspond to the 5' and 3' end sequences respectively as set forth in Example 1 using primers and having appropriate restriction sites and initiation/stop codons, if necessary. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to transform bacteria HB101, which are then plated onto

agar containing kanamycin for the purpose of confirming that the vector has the gene of interest properly inserted.

[01017] The amphotropic pA317 or GP+am12 packaging cells are grown in tissue culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is then added to the media and the packaging cells transduced with the vector. The packaging cells now produce infectious viral particles containing the gene (the packaging cells are now referred to as producer cells).

[01018] Fresh media is added to the transduced producer cells, and subsequently, the media is harvested from a 10 cm plate of confluent producer cells. The spent media, containing the infectious viral particles, is filtered through a millipore filter to remove detached producer cells and this media is then used to infect fibroblast cells. Media is removed from a sub-confluent plate of fibroblasts and quickly replaced with the media from the producer cells. This media is removed and replaced with fresh media. If the titer of virus is high, then virtually all fibroblasts will be infected and no selection is required. If the titer is very low, then it is necessary to use a retroviral vector that has a selectable marker, such as neo or his. Once the fibroblasts have been efficiently infected, the fibroblasts are analyzed to determine whether protein is produced.

[01019] The engineered fibroblasts are then transplanted onto the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads.

Example 17: Gene Therapy Using Endogenous Genes Corresponding To Polynucleotides of the Invention

[01020] Another method of gene therapy according to the present invention involves operably associating the endogenous polynucleotide sequence of the invention with a promoter via homologous recombination as described, for example, in U.S. Patent NO: 5,641,670, issued June 24, 1997; International Publication NO: WO 96/29411, published September 26, 1996; International Publication NO: WO 94/12650, published August 4, 1994; Koller et al., Proc. Natl. Acad. Sci. USA,

86:8932-8935 (1989); and Zijlstra et al., Nature, 342:435-438 (1989). This method involves the activation of a gene which is present in the target cells, but which is not expressed in the cells, or is expressed at a lower level than desired.

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[01021] Polynucleotide constructs are made which contain a promoter and targeting sequences, which are homologous to the 5' non-coding sequence of endogenous polynucleotide sequence, flanking the promoter. The targeting sequence will be sufficiently near the 5' end of the polynucleotide sequence so the promoter will be operably linked to the endogenous sequence upon homologous recombination. The promoter and the targeting sequences can be amplified using PCR. Preferably, the amplified promoter contains distinct restriction enzyme sites on the 5' and 3' ends. Preferably, the 3' end of the first targeting sequence contains the same restriction enzyme site as the 5' end of the amplified promoter and the 5' end of the second targeting sequence contains the same restriction site as the 3' end of the amplified promoter.

with the appropriate restriction enzymes and subsequently treated with calf intestinal phosphatase. The digested promoter and digested targeting sequences are added together in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The construct is size fractionated on an agarose gel, then purified by phenol extraction and ethanol precipitation.

[01023] In this Example, the polynucleotide constructs are administered as naked polynucleotides via electroporation. However, the polynucleotide constructs may also be administered with transfection-facilitating agents, such as liposomes, viral sequences, viral particles, precipitating agents, etc. Such methods of delivery are known in the art.

[01024] Once the cells are transfected, homologous recombination will take place which results in the promoter being operably linked to the endogenous polynucleotide sequence. This results in the expression of polynucleotide corresponding to the polynucleotide in the cell. Expression may be detected by immunological staining, or any other method known in the art.

is placed in DMEM + 10% fetal calf serum. Exponentially growing or early stationary phase fibroblasts are trypsinized and rinsed from the plastic surface with nutrient medium. An aliquot of the cell suspension is removed for counting, and the remaining cells are subjected to centrifugation. The supernatant is aspirated and the pellet is resuspended in 5 ml of electroporation buffer (20 mM HEPES pH 7.3, 137 mM NaCl. 5 mM KCl, 0.7 mM Na₂ HPO₄, 6 mM dextrose). The cells are recentrifuged, the supernatant aspirated, and the cells resuspended in electroporation buffer containing 1 mg/ml acetylated bovine serum albumin. The final cell suspension contains approximately 3X10⁶ cells/ml: Electroporation should be performed immediately following resuspension.

Plasmid DNA is prepared according to standard techniques. For example, to construct a plasmid for targeting to the locus corresponding to the polynucleotide of the invention, plasmid pUC18 (MBI Fermentas, Amherst, NY) is digested with HindIII. The CMV promoter is amplified by PCR with an XbaI site on the 5' end and a BamHI site on the 3' end. Two non-coding sequences are amplified via PCR: one non-coding sequence (fragment 1) is amplified with a HindIII site at the 5' end and an Xba site at the 3'end; the other non-coding sequence (fragment 2) is amplified with a BamHI site at the 5'end and a HindIII site at the 3'end. The CMV promoter and the fragments (1 and 2) are digested with the appropriate enzymes (CMV promoter - XbaI and BamHI; fragment 1 - XbaI; fragment 2 - BamHI) and ligated together. The resulting ligation product is digested with HindIII, and ligated with the HindIII-digested pUC18 plasmid.

[01027] Plasmid DNA is added to a sterile cuvette with a 0.4 cm' electrode gap (Bio-Rad). The final DNA concentration is generally at least 120 μg/ml. 0.5 ml of the cell suspension (containing approximately 1.5.X10⁶ cells) is then added to the cuvette, and the cell suspension and DNA solutions are gently mixed. Electroporation is performed with a Gene-Pulser apparatus (Bio-Rad). Capacitance and voltage are set at 960 μF and 250-300 V, respectively. As voltage increases, cell survival decreases, but the percentage of surviving cells that stably incorporate the introduced DNA into their genome increases dramatically. Given these parameters, a pulse time of approximately 14-20 mSec should be observed.

[01028] Electroporated cells are maintained at room temperature for approximately 5 min, and the contents of the cuvette are then gently removed with a sterile transfer pipette. The cells are added directly to 10 ml of prewarmed nutrient media (DMEM with 15% calf serum) in a 10 cm dish and incubated at 37 degree C. The following day, the media is aspirated and replaced with 10 ml of fresh media and incubated for a further 16-24 hours.

[01029] The engineered fibroblasts are then injected into the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads. The fibroblasts now produce the protein product. The fibroblasts can then be introduced into a patient as described above.

Example 18: Method of Treatment Using Gene Therapy - In Vivo

[01030] Another aspect of the present invention is using *in vivo* gene therapy methods to treat disorders, diseases and conditions. The gene therapy method relates to the introduction of naked nucleic acid (DNA, RNA, and antisense DNA or RNA) sequences into an animal to increase or decrease the expression of the polypeptide. The polynucleotide of the present invention may be operatively linked to (i.e., associated with) a promoter or any other genetic elements necessary for the expression of the polypeptide by the target tissue. Such gene therapy and delivery techniques and methods are known in the art, see, for example, WO90/11092, WO98/11779; U.S. Patent NO. 5693622, 5705151, 5580859; Tabata et al., Cardiovasc. Res. 35(3):470-479 (1997); Chao et al., Pharmacol. Res. 35(6):517-522 (1997); Wolff, Neuromuscul. Disord. 7(5):314-318 (1997); Schwartz et al., Gene Ther. 3(5):405-411 (1996); Tsurumi et al., Circulation 94(12):3281-3290 (1996) (incorporated herein by reference).

[01031] The polynucleotide constructs may be delivered by any method that delivers injectable materials to the cells of an animal, such as, injection into the interstitial space of tissues (heart, muscle, skin, lung, liver, intestine and the like). The polynucleotide constructs can be delivered in a pharmaceutically acceptable liquid or aqueous carrier.

[01032] The term "naked" polynucleotide, DNA or RNA, refers to sequences that 1188

are free from any delivery vehicle that acts to assist, promote, or facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. However, the polynucleotides of the present invention may also be delivered in liposome formulations (such as those taught in Felgner P.L. et al. (1995) Ann. NY Acad. Sci. 772:126-139 and Abdallah B. et al. (1995) Biol. Cell 85(1):1-7) which can be prepared by methods well known to those skilled in the art.

[01033] The polynucleotide vector constructs used in the gene therapy method are preferably constructs that will not integrate into the host genome nor will they contain sequences that allow for replication. Any strong promoter known to those skilled in the art can be used for driving the expression of DNA. Unlike other gene therapy techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired polypeptide for periods of up to six months.

The polynucleotide construct can be delivered to the interstitial space of [01034] tissues within an animal, including muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, blood, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, and connective tissue. Interstitial space of the tissues comprises the intercellular fluid, mucopolysaccharide matrix among the reticular fibers of organ tissues, elastic fibers in the walls of vessels or chambers, collagen fibers of fibrous tissues, or that same matrix within connective tissue ensheathing muscle cells or in the lacunae of bone. It is similarly the space occupied by the plasma of the circulation and the lymph fluid of the lymphatic channels. Delivery to the interstitial space of muscle tissue is preferred for the reasons discussed below. They may be conveniently delivered by injection into the tissues comprising these cells. They are preferably delivered to and expressed in persistent, non-dividing cells which are differentiated, although delivery and expression may be achieved in non-differentiated or less completely differentiated cells, such as, for example, stem cells of blood or skin fibroblasts. In vivo muscle cells are particularly competent in their ability to take up and express polynucleotides.

[01035] For the naked polynucleotide injection, an effective dosage amount of

DNA or RNA will be in the range of from about 0.05 g/kg body weight to about 50 mg/kg body weight. Preferably the dosage will be from about 0.005 mg/kg to about 20 mg/kg and more preferably from about 0.05 mg/kg to about 5 mg/kg. Of course, as the artisan of ordinary skill will appreciate, this dosage will vary according to the tissue site of injection. The appropriate and effective dosage of nucleic acid sequence can readily be determined by those of ordinary skill in the art and may depend on the condition being treated and the route of administration. The preferred route of administration is by the parenteral route of injection into the interstitial space of tissues. However, other parenteral routes may also be used, such as, inhalation of an aerosol formulation particularly for delivery to lungs or bronchial tissues, throat or mucous membranes of the nose. In addition, naked polynucleotide constructs can be delivered to arteries during angioplasty by the catheter used in the procedure.

[01036] The dose response effects of injected polynucleotide in muscle in vivo is determined as follows. Suitable template DNA for production of mRNA coding for polypeptide of the present invention is prepared in accordance with a standard recombinant DNA methodology. The template DNA, which may be either circular or linear, is either used as naked DNA or complexed with liposomes. The quadriceps muscles of mice are then injected with various amounts of the template DNA.

[01037] Five to six week old female and male Balb/C mice are anesthetized by intraperitoneal injection with 0.3 ml of 2.5% Avertin. A 1.5 cm incision is made on the anterior thigh, and the quadriceps muscle is directly visualized. The template DNA is injected in 0.1 ml of carrier in a 1 cc syringe through a 27 gauge needle over one minute, approximately 0.5 cm from the distal insertion site of the muscle into the knee and about 0.2 cm deep. A suture is placed over the injection site for future localization, and the skin is closed with stainless steel clips.

[01038] After an appropriate incubation time (e.g., 7 days) muscle extracts are prepared by excising the entire quadriceps. Every fifth 15 um cross-section of the individual quadriceps muscles is histochemically stained for protein expression. A time course for protein expression may be done in a similar fashion except that quadriceps from different mice are harvested at different times. Persistence of DNA in muscle following injection may be determined by Southern blot analysis after preparing total cellular DNA and HIRT supernatants from injected and control mice.

The results of the above experimentation in mice can be used to extrapolate proper dosages and other treatment parameters in humans and other animals using naked DNA.

Example 19: Transgenic Animals

[01039] The polypeptides of the invention can also be expressed in transgenic animals. Animals of any species, including, but not limited to, mice, rats, rabbits, hamsters, guinea pigs, pigs, micro-pigs, goats, sheep, cows and non-human primates, e.g., baboons, monkeys, and chimpanzees may be used to generate transgenic animals. In a specific embodiment, techniques described herein or otherwise known in the art, are used to express polypeptides of the invention in humans, as part of a gene therapy protocol.

Any technique known in the art may be used to introduce the transgene [01040] (i.e., polynucleotides of the invention) into animals to produce the founder lines of transgenic animals. Such techniques include, but are not limited to, pronuclear microinjection (Paterson et al., Appl. Microbiol. Biotechnol. 40:691-698 (1994); Carver et al., Biotechnology (NY) 11:1263-1270 (1993); Wright et al., Biotechnology (NY) 9:830-834 (1991); and Hoppe et al., U.S. Pat. No. 4,873,191 (1989)); retrovirus mediated gene transfer into germ lines (Van der Putten et al., Proc. Natl. Acad. Sci., USA 82:6148-6152 (1985)), blastocysts or embryos; gene targeting in embryonic stem cells (Thompson et al., Cell 56:313-321 (1989)); electroporation of cells or embryos (Lo, 1983, Mol Cell. Biol. 3:1803-1814 (1983)); introduction of the polynucleotides of the invention using a gene gun (see, e.g., Ulmer et al., Science 259:1745 (1993); introducing nucleic acid constructs into embryonic pleuripotent stem cells and transferring the stem cells back into the blastocyst; and sperm-mediated gene transfer (Lavitrano et al., Cell 57:717-723 (1989); etc. For a review of such techniques, see Gordon, "Transgenic Animals," Intl. Rev. Cytol. 115:171-229 (1989), which is incorporated by reference herein in its entirety.

[01041] Any technique known in the art may be used to produce transgenic clones containing polynucleotides of the invention, for example, nuclear transfer into enucleated oocytes of nuclei from cultured embryonic, fetal, or adult cells induced to

quiescence (Campell et al., Nature 380:64-66 (1996); Wilmut et al., Nature 385:810-813 (1997)).

The present invention provides for transgenic animals that carry the [01042] transgene in all their cells, as well as animals which carry the transgene in some, but not all their cells, i.e., mosaic animals or chimeric. The transgene may be integrated as a single transgene or as multiple copies such as in concatamers, e.g., head-to-head tandems or head-to-tail tandems. The transgene may also be selectively introduced into and activated in a particular cell type by following, for example, the teaching of Lasko et al. (Lasko et al., Proc. Natl. Acad. Sci. USA 89:6232-6236 (1992)). The regulatory sequences required for such a cell-type specific activation will depend upon the particular cell type of interest, and will be apparent to those of skill in the art. When it is desired that the polynucleotide transgene be integrated into the chromosomal site of the endogenous gene, gene targeting is preferred. Briefly, when such a technique is to be utilized, vectors containing some nucleotide sequences homologous to the endogenous gene are designed for the purpose of integrating, via homologous recombination with chromosomal sequences, into and disrupting the function of the nucleotide sequence of the endogenous gene. The transgene may also be selectively introduced into a particular cell type, thus inactivating the endogenous gene in only that cell type, by following, for example, the teaching of Gu et al. (Gu et al., Science 265:103-106 (1994)). The regulatory sequences required for such a celltype specific inactivation will depend upon the particular cell type of interest, and will be apparent to those of skill in the art.

[01043] Once transgenic animals have been generated, the expression of the recombinant gene may be assayed utilizing standard techniques. Initial screening may be accomplished by Southern blot analysis or PCR techniques to analyze animal tissues to verify that integration of the transgene has taken place. The level of mRNA expression of the transgene in the tissues of the transgenic animals may also be assessed using techniques which include, but are not limited to, Northern blot analysis of tissue samples obtained from the animal, in situ hybridization analysis, and reverse transcriptase-PCR (rt-PCR). Samples of transgenic gene-expressing tissue may also be evaluated immunocytochemically or immunohistochemically using antibodies specific for the transgene product.

Once the founder animals are produced, they may be bred, inbred, outbred, or crossbred to produce colonies of the particular animal. Examples of such breeding strategies include, but are not limited to: outbreeding of founder animals with more than one integration site in order to establish separate lines; inbreeding of separate lines in order to produce compound transgenics that express the transgene at higher levels because of the effects of additive expression of each transgene; crossing of heterozygous transgenic animals to produce animals homozygous for a given integration site in order to both augment expression and eliminate the need for screening of animals by DNA analysis; crossing of separate homozygous lines to produce compound heterozygous or homozygous lines; and breeding to place the transgene on a distinct background that is appropriate for an experimental model of interest.

[01045] Transgenic animals of the invention have uses which include, but are not limited to, animal model systems useful in elaborating the biological function of polypeptides of the present invention, studying conditions and/or disorders associated with aberrant expression, and in screening for compounds effective in ameliorating such conditions and/or disorders.

Example 20: Knock-Out Animals

"knocking out" the gene and/or its promoter using targeted homologous recombination. (See e.g., Smithies et al., Nature 317:230-234 (1985); Thomas & Capecchi, Cell 51:503-512 (1987); Thompson et al., Cell 5:313-321 (1989); each of which is incorporated by reference herein in its entirety.) For example, a mutant, non-functional polynucleotide of the invention (or a completely unrelated DNA sequence) flanked by DNA homologous to the endogenous polynucleotide sequence (either the coding regions or regulatory regions of the gene) can be used, with or without a selectable marker and/or a negative selectable marker, to transfect cells that express polypeptides of the invention in vivo. In another embodiment, techniques known in the art are used to generate knockouts in cells that contain, but do not express the gene of interest. Insertion of the DNA construct, via targeted homologous recombination,

results in inactivation of the targeted gene. Such approaches are particularly suited in research and agricultural fields where modifications to embryonic stem cells can be used to generate animal offspring with an inactive targeted gene (e.g., see Thomas & Capecchi 1987 and Thompson 1989, *supra*). However this approach can be routinely adapted for use in humans provided the recombinant DNA constructs are directly administered or targeted to the required site *in vivo* using appropriate viral vectors that will be apparent to those of skill in the art.

[01047] In further embodiments of the invention, cells that are genetically engineered to express the polypeptides of the invention, or alternatively, that are genetically engineered not to express the polypeptides of the invention (e.g., knockouts) are administered to a patient in vivo. Such cells may be obtained from the patient (i.e., animal, including human) or an MHC compatible donor and can include, but are not limited to fibroblasts, bone marrow cells, blood cells (e.g., lymphocytes), adipocytes, muscle cells, endothelial cells etc. The cells are genetically engineered in vitro using recombinant DNA techniques to introduce the coding sequence of polypeptides of the invention into the cells, or alternatively, to disrupt the coding sequence and/or endogenous regulatory sequence associated with the polypeptides of the invention, e.g., by transduction (using viral vectors, and preferably vectors that integrate the transgene into the cell genome) or transfection procedures, including, but not limited to, the use of plasmids, cosmids, YACs, naked DNA, electroporation, liposomes, etc. The coding sequence of the polypeptides of the invention can be placed under the control of a strong constitutive or inducible promoter or promoter/enhancer to achieve expression, and preferably secretion, of the polypeptides of the invention. The engineered cells which express and preferably secrete the polypeptides of the invention can be introduced into the patient systemically, e.g., in the circulation, or intraperitoneally.

[01048] Alternatively, the cells can be incorporated into a matrix and implanted in the body, e.g., genetically engineered fibroblasts can be implanted as part of a skin graft; genetically engineered endothelial cells can be implanted as part of a lymphatic or vascular graft. (See, for example, Anderson et al. U.S. Patent No. 5,399,349; and Mulligan & Wilson, U.S. Patent No. 5,460,959 each of which is incorporated by reference herein in its entirety).

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